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(54) Title: METHODS OF DIAGNOSIS OF LUNG CANCER, COMPOSITIONS AND METHODS OF SCREENING FOR MODULATORS OF LUNG CANCER

(57) Abstract: Described herein are methods and compositions that can be used for diagnosis and treatment of lung cancer and similar pathologies. Also described herein are methods that can be used to identify modulators of lung cancer and similar pathologies.

METHODS OF DIAGNOSIS OF LUNG CANCER, COMPOSITIONS AND METHODS OF SCREENING FOR MODULATORS OF LUNG CANCER

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CROSS-REFERENCES TO RELATED APPLICATIONS

This application is related to USSN 60/284,770, filed April 18, 2001; USSN 60/290,492, filed May 10, 2001; USSN 60/334,370, filed November 29, 2001; USSN 60/339,245, filed November 9, 2001; USSN 60/350,666, filed November 13, 2001; and USSN 60/xxx,xxx, filed April 12, 2002 (Docket OMNI-002P); each of which is incorporated herein by reference in its entirety.

FIELD OF THE INVENTION

The invention relates to the identification of nucleic acid and protein expression profiles and nucleic acids, products, and antibodies thereto that are involved in lung cancer; and to the use of such expression profiles and compositions in diagnosis and therapy of lung cancer. The invention further relates to methods for identifying and using agents and/or targets that inhibit lung cancer or related conditions.

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BACKGROUND OF THE INVENTION

Lung cancer is the second most commonly occurring cancer in the United States and is the leading cause of cancer-related death. It is estimated that there are over 160,000 new cases of lung cancer in the United States every year. Of those who are diagnosed with lung cancer, 86 percent will die within five years. Lung cancer is the most common visceral cancer in men and accounts for nearly one third of all cancer deaths in both men and women. In fact, lung cancer accounts for 7% of all deaths, due to any cause, in both men and women.

Smoking is the primary cause of lung cancer, with more than 80% of lung cancers resulting from smoking. About 400 to 500 separate gaseous substances are present in the smoke of a non-filter cigarette. The most noteworthy substances include nitrogen oxides, hydrogen cyanide, formaldehyde, benzene, and toluene. The particles present in cigarette smoke contain at least 3,500 individual compounds such as nicotine, tobacco alkaloids (nornicotine, anatabine, anabasine), polycyclic aromatic hydrocarbons (e.g., benzo(a)pyrene, B(a)P), naphthalenes, aromatic amines, phenols, and tobacco-specific nitrosamines.

Tobacco-specific nitrosamines are formed during tobacco curing and processing, and are suspected of causing lung cancer in humans. In rodent studies, regardless of the where or how it is applied, the tobacco-specific nitrosamine known as NNK produces lung adenomas and lung adenocarcinomas. The tobacco-specific nitrosamine known as NNAL also produces lung adenocarcinomas in rodents.

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Many of the chemicals found in cigarette smoke also affect the nonsmoker inhaling "secondhand" or sidestream smoke. Indeed, the smoke inhaled by non-smokers has a chemical composition similar to the smoke inhaled by smokers, but, importantly, the concentrations of the carcinogenic tobacco-specific nitrosamines are present in higher concentrations in second hand smoke. For this and other reasons, "passive smoking" is an important cause of lung cancer, causing as many as 3,000 lung cancer deaths in nonsmokers each year.

In addition to smoking, other factors thought to be causes of lung cancer include onthe-job exposure to carcinogens such as asbestos and uranium, exposure to chemical hazards such as radon, polycyclic aromatic hydrocarbons, chromium, nickel, and inorganic arsenic, genetic factors, and diet.

Histological classification of various lung cancers define the types of cancer that begin in the lung. See, e.g., Travis, et al. (1999) <u>Histological Typing of Lung and Pleural Tumours</u> (International Histological Classification of Tumours, No 1. Four major cell types make up more than 88% of all primary lung neoplasms. These are: squamous or epidermoid carcinoma, small cell (also called oat cell) carcinoma, adenocarcinoma, and large cell (also called large cell anaplastic) carcinoma. The remainder include undifferentiated carcinomas, carcinoids, bronchial gland tumors, and other rarer types. The various cell types have different natural histories and responses to therapy, and, thus, a correct histologic diagnosis is the first step of effective treatment.

Small cell lung cancer (SCLC) accounts for 18-25% of all lung cancers, and occurs less frequently than non-small cell lung cancers, and generally spread to distant organs more rapidly than non-small cell lung cancer. In general, at the time of presentation small cell lung cancers have already spread beyond the beyond the bounds where surgery and curative intent can be undertaken. Hoever, if identified early enough, these cancers are often responsive to chemotherapy and thoracic radiation treatment.

Non-small cell lung cancers (NSCLC) are the more frequently occurring form of lung cancer. They comprise squamous cell carcinoma, adenocarcinoma, and large cell carcinoma

and account for more than 75% of all lung cancers. Non-small cell tumors that are localized at the time of presentation can sometimes be cured with surgery and/or radiotherapy, but usually are not identified until significant metastasis has occurred, which are typically not very responsive to surgical, chemotherapy, or radiation treatment..

The screening of asymptomatic persons at high risk for lung cancer has often proven ineffective. In general, only 5 to 15 percent of lung cancer patients have their disease detected while they are asymptomatic. Of course, early detection and treatment are critical factors in the fight against lung cancer. The average survival rate is 49% for those whose cancer is detected early, before the cancer has spread from the lung. Lung cancer often spreads outside of the lung, and it may have spread to the bones or brain by the time it is diagnosed. While the prognosis may be better for lung cancers that are detected early, because of the lack of effective curative treatments, early detection does not necessarily alter the total death rate from lung cancer.

Thus, methods for diagnosis and prognosis of lung cancer and effective treatment of lung cancer would be desirable. Accordingly, provided herein are methods that can be used in diagnosis and prognosis of lung cancer. Further provided are methods that can be used to screen candidate therapeutic agents for the ability to modulate, e.g., treat, lung cancer. Additionally, provided herein are molecular targets and compositions for therapeutic intervention in lung disease and other metastatic cancers.

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SUMMARY OF THE INVENTION

The present invention provides nucleotide sequences of genes that are up- and down-regulated in lung cancer cells. Such genes are useful for diagnostic purposes, and also as targets for screening for therapeutic compounds that modulate lung cancer, such as antibodies. The methods of detecting nucleic acids of the invention or their encoded proteins can be used for a number of purposes. Examples include early detection of lung cancers, monitoring and early detection of relapse following treatment of lung cancers, monitoring response to therapy of lung cancers, determining prognosis of lung cancers, directing therapy of lung cancers, selecting patients for postoperative chemotherapy or radiation therapy, selecting therapy, determining tumor prognosis, treatment, or response to treatment, and early detection of precancerous lesions of the lung. Examples of benign or precancerous lesions include: atelectasis, emphysema, brochitis, chronic obstructive pulmonary disease, fibrosis, hypersensitivity pneumonitis (HP), interstitial pulmonary fibrosis (IPF), asthma, and

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bronchiectasis. Other aspects of the invention will become apparent to the skilled artisan by
the following description of the invention.

In one aspect, the present invention provides a method of detecting a lung cancer-associated transcript in a cell from a patient, the method comprising contacting a biological sample from the patient with a polynucleotide that selectively hybridizes to a sequence at least 80% identical to a sequence as shown in Tables 1A-16. Alternatively, the sample may be contacted with a specific binding reagent, e.g., antibody.

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In one embodiment, the polynucleotide selectively hybridizes to a sequence at least 95% identical to a sequence as shown in Tables 1A-16. In another embodiment, the polynucleotide comprises a sequence as shown in Tables 1A-16.

In one embodiment, the biological sample is a tissue sample, or a body fluid. In another embodiment, the biological sample comprises isolated nucleic acids, e.g., mRNA.

In one embodiment, the polynucleotide is labeled, e.g., with a fluorescent label. In one embodiment, the polynucleotide is immobilized on a solid surface. In one embodiment, the patient is undergoing a therapeutic regimen to treat lung cancer. In another embodiment, the patient is suspected of having lung cancer. In one embodiment, the patient is a primate, e.g., a human.

In one embodiment, the method further comprises the step of amplifying nucleic acids before the step of contacting the biological sample with the polynucleotide.

In another aspect, the present invention provides a method of monitoring the efficacy of a therapeutic treatment of lung cancer, the method comprising the steps of: (i) providing a biological sample from a patient undergoing the therapeutic treatment; and (ii) determining the level of a lung cancer-associated transcript in the biological sample by contacting the biological sample with a polynucleotide that selectively hybridizes to a sequence at least 80% identical to a sequence as shown in Tables 1A-16, thereby monitoring the efficacy of the therapy. Or the sample may be evaluated for protein, e.g., contacting the sample with an antibody.

In one embodiment, the method further comprises the step of: (iii) comparing the level of the lung cancer-associated transcript to a level of the lung cancer-associated transcript in a biological sample from the patient prior to, or earlier in, the therapeutic treatment. Or the sample may be evaluated for comparison of protein.

In another aspect, the present invention provides a method of monitoring the efficacy of a therapeutic treatment of lung cancer, the method comprising the steps of: (i) providing a

biological sample from a patient undergoing the therapeutic treatment; and (ii) determining the level of a lung cancer-associated antibody in the biological sample by contacting the biological sample with a polypeptide encoded by a polynucleotide that selectively hybridizes to a sequence at least 80% identical to a sequence as shown in Tables 1A-16, wherein the polypeptide specifically binds to the lung cancer-associated antibody, thereby monitoring the efficacy of the therapy.

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In one embodiment, the method further comprises the step of: (iii) comparing the level of the lung cancer-associated antibody to a level of the lung cancer-associated antibody in a biological sample from the patient prior to, or earlier in, the therapeutic treatment.

In another aspect, the present invention provides a method of monitoring the efficacy of a therapeutic treatment of lung cancer, the method comprising the steps of: (i) providing a biological sample from a patient undergoing the therapeutic treatment; and (ii) determining the level of a lung cancer-associated polypeptide in the biological sample by contacting the biological sample with an antibody, wherein the antibody specifically binds to a polypeptide encoded by a polynucleotide that selectively hybridizes to a sequence at least 80% identical to a sequence as shown in Tables 1A-16, thereby monitoring the efficacy of the therapy.

In one embodiment, the method further comprises the step of: (iii) comparing the level of the lung cancer-associated polypeptide to a level of the lung cancer-associated polypeptide in a biological sample from the patient prior to, or earlier in, the therapeutic treatment. In one aspect, the present invention provides an isolated nucleic acid molecule consisting of a polynucleotide sequence as shown in Tables 1A-16. In one embodiment, an expression vector or cell comprises the isolated nucleic acid. In one aspect, the present invention provides an isolated polypeptide which is encoded by a nucleic acid molecule having polynucleotide sequence as shown in Tables 1A-16.

In another aspect, the present invention provides an antibody that specifically binds to an isolated polypeptide which is encoded by a nucleic acid molecule having polynucleotide sequence as shown in Tables 1A-16. In one embodiment, the antibody is conjugated to an effector component, e.g., a fluorescent label, a radioisotope or a cytotoxic chemical. In one embodiment, the antibody is an antibody fragment. In another embodiment, the antibody is humanized.

In one aspect, the present invention provides a method of detecting lung cancer in a a patient, the method comprising contacting a biological sample from the patient with an antibody or protein as described herein.

In another aspect, the present invention provides a method of detecting antibodies specific to a lung cancer gene in a patient, the method comprising contacting a biological sample from the patient with a polypeptide encoded by a nucleic acid comprises a sequence from Tables 1A-16.

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In another aspect, the present invention provides a method for identifying a compound that modulates a lung cancer-associated polypeptide, the method comprising the steps of: (i) contacting the compound with a lung cancer-associated polypeptide, the polypeptide encoded by a polynucleotide that selectively hybridizes to a sequence at least 80% identical to a sequence as shown in Tables 1A-16; and (ii) determining the functional effect of the compound upon the polypeptide.

In one embodiment, the functional effect is a physical effect, an enzymatic effect, or a chemical effect. In one embodiment, the polypeptide is expressed in a eukaryotic host cell or cell membrane. In another embodiment, the polypeptide is recombinant. In one embodiment, the functional effect is determined by measuring ligand binding to the polypeptide.

In another aspect, the present invention provides a method of inhibiting proliferation or another critical process of a lung cancer-associated cell to treat lung cancer in a patient, the method comprising the step of administering to the subject a therapeutically effective amount of a compound identified as described herein. In one embodiment, the compound is an antibody.

In another aspect, the present invention provides a drug screening assay comprising the steps of: (i) administering a test compound to a mammal having lung cancer or a cell isolated therefrom; (ii) comparing the level of gene expression of a polynucleotide that selectively hybridizes to a sequence at least 80% identical to a sequence as shown in Tables 1A-16 in a treated cell or mammal with the level of gene expression of the polynucleotide in a control cell or mammal, wherein a test compound that modulates the level of expression of the polynucleotide is a candidate for the treatment of lung cancer.

In one embodiment, the control is a mammal with lung cancer or a cell therefrom that has not been treated with the test compound. In another embodiment, the control is a normal cell or mammal, or a non-malignant lung disease.

In another aspect, the present invention provides a method for treating a mammal having lung cancer comprising administering a compound identified by the assay described herein.

In another aspect, the present invention provides a pharmaceutical composition for treating a mammal having lung cancer, the composition comprising a compound identified by the assay described herein and a physiologically acceptable excipient.

DETAILED DESCRIPTION OF THE INVENTION

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In accordance with the objects outlined above, the present invention provides novel methods for diagnosis and treatment of lung disease or cancer, as well as methods for screening for compositions which modulate lung cancer. "Treatment, monitoring, detection or modulation of lung disease or cancer" includes treatment, monitoring, detection, or modulation of lung disease in those patients who have lung disease (whether malignant or non-malignant, e.g., emphysema, bronchitis, or fibrosis) as well as patients with lung cancers in which gene expression from a gene in Tables 1A-16 is increased or decreased, indicating that the subject is more likely to have disease. In particular, while these targets are identified primarily from lung cancer samples, these same targets are likely to be similarly found in analyses of other medical conditions. These other conditions may result from similar pathological processes which affect similar tissues, e.g., lung cancer, small cell lung carcinoma (oat cell carcinoma), non-small cell carcinomas (e.g., squamous cell carcinoma, adenocarcinoma, large cell lung carcinoma, carcinoid, granulomatous), fibrosis (idiopathic pulmonary fibrosis (IPF), hypersensitivity pneumonitis (HP), interstitial pneumonitis, nonspecific idiopathic pneumonitis (NSIP)), chronic obstructive pulmonary disease (COPD, e.g., emphysema, chronic bronchitis), asthma, bronchiectasis, and esophageal cancer. See, e.g., the NCI webpage and USSN 60/347,349 and USSN 60/xxx,xxx (docket LFBR-001-1P, filed March 29, 2002), each of which is incorporated herein by reference. The treatment may be of lung cancer or related condition itself, or treatment of metastasis.

In particular, identification of markers selectively expressed on these cancers allows for use of that expression in diagnostic, prognostic, or therapeutic methods. As such, the invention defines various compositions, e.g., nucleic acids, polypeptides, antibodies, and small molecule agonists/antagonists, which will be useful to selectively identify those markers. For example, therapeutic methods may take the form of protein therapeutics which use the marker expression for selective localization or modulation of function (for those markers which have a causative disease effect), for vaccines, identification of binding partners, or antagonism, e.g., using antisense or RNAi. The markers may be useful for molecular characterization of subsets of lung diseases, which subsets may actually require

very different treatments. Moreover, the markers may also be important in related diseases to the specific cancers, e.g., which affect similar tissues in non-malignant diseases, or have similar mechanisms of induction/maintenance. Metastatic processes or characteristics may also be targeted. Diagnostic and prognostic uses are made available, e.g., to subset related but distinct diseases, or to determine treatment strategy. The detection methods may be based upon nucleic acid, e.g., PCR or hybridization techniques, or protein, e.g., ELISA, imaging, IHC, etc. The diagnosis may be qualitative or quantitative, and may detect increases or decreases in expression levels.

Tables 1A-16 provide unigene cluster identification numbers for the nucleotide sequence of genes that exhibit increased or decreased expression in lung cancer samples. The tables also provide an exemplar accession number that provides a nucleotide sequence that is part of the unigene cluster. In Table 1A, genes marked as "target 1" or "target 2" are particularly useful as therapeutic targets. Genes marked as "target 3" are particularly useful as diagnostic markers. Genes marked as "chron" are upregulated in chronically diseased lung (e.g., emphysema, bronchitis, fibrosis) relative to lung tumors and normal tissue. In certain analyses, the ratio for the "chron" category was determined using the 70th percentile of chronically diseases lung samples divided by the 90th percentile of lung tumor samples divided by the 90th percentile of lung tumor samples divided by the 90th percentile of normal lung samples.

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Definitions

The term "lung cancer protein" or "lung cancer polynucleotide" or "lung cancerassociated transcript" refers to nucleic acid and polypeptide polymorphic variants, alleles, mutants, and interspecies homologs that: (1) have a nucleotide sequence that has greater than about 60% nucleotide sequence identity, 65%, 70%, 75%, 80%, 85%, 90%, preferably 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, or 99% or greater nucleotide sequence identity, preferably over a region of over a region of at least about 25, 50, 100, 200, 500, 1000, or more nucleotides, to a nucleotide sequence of or associated with a unigene cluster of Tables 1A-16; (2) bind to antibodies, e.g., polyclonal antibodies, raised against an immunogen comprising an amino acid sequence encoded by a nucleotide sequence of or associated with a unigene cluster of Tables 1A-16, and conservatively modified variants thereof; (3) specifically hybridize under stringent hybridization conditions to a nucleic acid sequence, or the complement thereof of Tables 1A-16 and conservatively modified variants thereof; or (4)

have an amino acid sequence that has greater than about 60% amino acid sequence identity, 65%, 70%, 75%, 80%, 85%, 90%, preferably 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, or 99% or greater amino sequence identity, preferably over a region of over a region of at least about 25, 50, 100, 200, 500, 1000, or more amino acid, to an amino acid sequence encoded by a nucleotide sequence of or associated with a unigene cluster of Tables 1A-16. A polynucleotide or polypeptide sequence is typically from a mammal including, but not limited to, primate, e.g., human; rodent, e.g., rat, mouse, hamster; cow, pig, horse, sheep, or other mammal. A "lung cancer polypeptide" and a "lung cancer polynucleotide," include both naturally occurring or recombinant forms.

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A "full length" lung cancer protein or nucleic acid refers to a lung cancer polypeptide or polynucleotide sequence, or a variant thereof, that contains the elements normally contained in one or more naturally occurring, wild type lung cancer polynucleotide or polypeptide sequences. The "full length" may be prior to, or after, various stages of post-translational processing or splicing, including alternative splicing.

"Biological sample" as used herein is a sample of biological tissue or fluid that contains nucleic acids or polypeptides, e.g., of a lung cancer protein, polynucleotide, or transcript. Such samples include, but are not limited to, tissue isolated from primates, e.g., humans, or rodents, e.g., mice, and rats. Biological samples may also include sections of tissues such as biopsy and autopsy samples, frozen sections taken for histologic purposes, archival materials, blood, plasma, serum, sputum, stool, tears, mucus, hair, skin, etc. Biological samples also include explants and primary and/or transformed cell cultures derived from patient tissues. A biological sample is typically obtained from a eukaryotic organism, most preferably a mammal such as a primate, e.g., chimpanzee or human; cow; dog; cat; a rodent, e.g., guinea pig, rat, mouse; rabbit; or other mammal; or a bird; reptile; fish. Livestock and domestic animals are of interest.

"Providing a biological sample" means to obtain a biological sample for use in methods described in this invention. Most often, this will be done by removing a sample of cells from an animal, but can also be accomplished by using previously isolated cells (e.g., isolated by another person, at another time, and/or for another purpose), or by performing the methods of the invention in vivo. Archival tissues or materials, having treatment or outcome history, will be particularly useful.

The terms "identical" or percent "identity," in the context of two or more nucleic acids or polypeptide sequences, refer to two or more sequences or subsequences that are the

same or have a specified percentage of amino acid residues or nucleotides that are the same (e.g., about 60% identity, preferably 70%, 75%, 80%, 85%, 90%, 91%, 92%, 93%, 94%, 95%, 96%, 97%, 98%, 99%, or higher identity over a specified region, when compared and aligned for maximum correspondence over a comparison window or designated region) as measured using, e.g., a BLAST or BLAST 2.0 sequence comparison algorithms with default parameters described below, or by manual alignment and visual inspection (see, e.g., NCBI web site http://www.ncbi.nlm.nih.gov/BLAST/ or the like). Such sequences are then said to be "substantially identical." This definition also refers to, or may be applied to, the complement of a test sequence. The definition also includes sequences that have deletions and/or insertions, substitutions, and naturally occurring, e.g., polymorphic or allelic variants, and man-made variants. As described below, the preferred algorithms can account for gaps and the like. Preferably, identity exists over a region that is at least about 25 amino acids or nucleotides in length, or more preferably over a region that is 50-100 amino acids or nucleotides in length.

For sequence comparison, typically one sequence acts as a reference sequence, to which test sequences are compared. When using a sequence comparison algorithm, test and reference sequences are entered into a computer, subsequence coordinates are designated, if necessary, and sequence algorithm program parameters are designated. Preferably, default program parameters can be used, or alternative parameters can be designated. The sequence comparison algorithm then calculates the percent sequence identities for the test sequences relative to the reference sequence, based on the program parameters.

A "comparison window", as used herein, includes reference to a segment of contiguous positions selected from the group consisting typically of from 20 to 600, usually about 50 to about 200, more usually about 100 to about 150 in which a sequence may be compared to a reference sequence of the same number of contiguous positions after the two sequences are optimally aligned. Methods of alignment of sequences for comparison are well-known in the art. Optimal alignment of sequences for comparison can be conducted, e.g., by the local homology algorithm of Smith and Waterman (1981) Adv. Appl. Math. 2:482, by the homology alignment algorithm of Needleman and Wunsch (1970) J. Mol. Biol. 48:443, by the search for similarity method of Pearson and Lipman (1988) Proc. Nat'l. Acad. Sci. USA 85:2444, by computerized implementations of these algorithms (GAP, BESTFIT, FASTA, and TFASTA in the Wisconsin Genetics Software Package, Genetics Computer

WO 02/086443 PCT/US02/12476 Group, 575 Science Dr., Madison, WI), or by manual alignment and visual inspection (see, e.g., Ausubel, et al. (eds. 1995 and supplements) <u>Current Protocols in Molecular Biology</u>.

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Preferred examples of algorithms that are suitable for determining percent sequence identity and sequence similarity include the BLAST and BLAST 2.0 algorithms, which are described in Altschul, et al. (1977) Nuc. Acids Res. 25:3389-3402 and Altschul, et al. (1990) J. Mol. Biol. 215:403-410. BLAST and BLAST 2.0 are used, with the parameters described herein, to determine percent sequence identity for the nucleic acids and proteins of the invention. Software for performing BLAST analyses is publicly available through the National Center for Biotechnology Information (http://www.ncbi.nlm.nih.gov/). This algorithm involves first identifying high scoring sequence pairs (HSPs) by identifying short words of length W in the query sequence, which either match or satisfy some positive-valued threshold score T when aligned with a word of the same length in a database sequence. T is referred to as the neighborhood word score threshold (Altschul, et al., supra). These initial neighborhood word hits act as seeds for initiating searches to find longer HSPs containing them. The word hits are extended in both directions along each sequence for as far as the cumulative alignment score can be increased. Cumulative scores are calculated using, e.g., for nucleotide sequences, the parameters M (reward score for a pair of matching residues; always > 0) and N (penalty score for mismatching residues; always < 0). For amino acid sequences, a scoring matrix is used to calculate the cumulative score. Extension of the word hits in each direction are halted when: the cumulative alignment score falls off by the quantity X from its maximum achieved value; the cumulative score goes to zero or below, due to the accumulation of one or more negative-scoring residue alignments; or the end of either sequence is reached. The BLAST algorithm parameters W, T, and X determine the sensitivity and speed of the alignment. The BLASTN program (for nucleotide sequences) uses as defaults a wordlength (W) of 11, an expectation (E) of 10, M=5, N=-4 and a comparison of both strands. For amino acid sequences, the BLASTP program uses as defaults a wordlength of 3, and expectation (E) of 10, and the BLOSUM62 scoring matrix (see Henikoff and Henikoff (1989) Proc. Natl. Acad. Sci. USA 89:10915) alignments (B) of 50, expectation (E) of 10, M=5, N=-4, and a comparison of both strands.

The BLAST algorithm also performs a statistical analysis of the similarity between two sequences (see, e.g., Karlin and Altschul (1993) <u>Proc. Nat'l. Acad. Sci. USA</u> 90:5873-5787). One measure of similarity provided by the BLAST algorithm is the smallest sum probability (P(N)), which provides an indication of the probability by which a match between

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is considered similar to a reference sequence if the smallest sum probability in a comparison of the test nucleic acid to the reference nucleic acid is less than about 0.2, more preferably less than about 0.01, and most preferably less than about 0.001. Log values may be negative large numbers, e.g., 5, 10, 20, 30, 40, 40, 70, 90, 110, 150, 170, etc.

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An indication that two nucleic acid sequences are substantially identical is that the polypeptide encoded by the first nucleic acid is immunologically cross reactive with the antibodies raised against the polypeptide encoded by the second nucleic acid. Thus, a polypeptide is typically substantially identical to a second polypeptide, e.g., where the two peptides differ only by conservative substitutions. Another indication that two nucleic acid sequences are substantially identical is that the two molecules or their complements hybridize to each other under stringent conditions. Yet another indication that two nucleic acid sequences are substantially identical is that the same primers can be used to amplify the sequences.

A "host cell" is a naturally occurring cell or a transformed cell that contains an expression vector and supports the replication or expression of the expression vector. Host cells may be cultured cells, explants, cells *in vivo*, and the like. Host cells may be prokaryotic cells such as *E. coli*, or eukaryotic cells such as yeast, insect, amphibian, or mammalian cells such as CHO, HeLa, and the like (see, e.g., the American Type Culture Collection catalog or web site, www.atcc.org).

The terms "isolated," "purified," or "biologically pure" refer to material that is substantially or essentially free from components that normally accompany it as found in its native state. Purity and homogeneity are typically determined using analytical chemistry techniques such as polyacrylamide gel electrophoresis or high performance liquid chromatography. A protein or nucleic acid that is the predominant species present in a preparation is substantially purified. In particular, an isolated nucleic acid is separated from some open reading frames that naturally flank the gene and encode proteins other than protein encoded by the gene. The term "purified" in some embodiments denotes that a nucleic acid or protein gives rise to essentially one band in an electrophoretic gel. Preferably, it means that the nucleic acid or protein is at least about 85% pure, more preferably at least 95% pure, and most preferably at least 99% pure. "Purify" or "purification" in other embodiments means removing at least one contaminant or component from the composition to be purified.

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In this sense, purification does not require that the purified compound be homogeneous, e.g., 100% pure.

The terms "polypeptide," "peptide" and "protein" are used interchangeably herein to refer to a polymer of amino acid residues. The terms apply to amino acid polymers in which one or more amino acid residue is an artificial chemical mimetic of a corresponding naturally occurring amino acid, as well as to naturally occurring amino acid polymers, those containing modified residues, and non-naturally occurring amino acid polymer.

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The term "amino acid" refers to naturally occurring and synthetic amino acids, as well as amino acid analogs and amino acid mimetics that function similarly to the naturally occurring amino acids. Naturally occurring amino acids are those encoded by the genetic code, as well as those amino acids that are later modified, e.g., hydroxyproline, γ-carboxyglutamate, and O-phosphoserine. Amino acid analogs refer to compounds that have the same basic chemical structure as a naturally occurring amino acid, e.g., an α carbon that is bound to a hydrogen, a carboxyl group, an amino group, and an R group, e.g., homoserine, norleucine, methionine sulfoxide, methionine methyl sulfonium. Such analogs may have modified R groups (e.g., norleucine) or modified peptide backbones, but retain some basic chemical structure as a naturally occurring amino acid. Amino acid mimetics refer to chemical compounds that have a structure that is different from the general chemical structure of an amino acid, but that function similarly to another amino acid.

Amino acids may be referred to herein by either their commonly known three letter symbols or by the one-letter symbols recommended by the IUPAC-IUB Biochemical Nomenclature Commission. Nucleotides, likewise, may be referred to by their commonly accepted single-letter codes.

"Conservatively modified variants" applies to both amino acid and nucleic acid sequences. With respect to particular nucleic acid sequences, conservatively modified variants refers to those nucleic acids which encode identical or essentially identical amino acid sequences, or where the nucleic acid does not encode an amino acid sequence, to essentially identical or associated, e.g., naturally contiguous, sequences. Because of the degeneracy of the genetic code, a large number of functionally identical nucleic acids encode most proteins. For instance, the codons GCA, GCC, GCG, and GCU each encode the amino acid alanine. Thus, at each position where an alanine is specified by a codon, the codon can be altered to another of the corresponding codons described without altering the encoded polypeptide. Such nucleic acid variations are "silent variations," which are one species of

WO 02/086443 PCT/US02/12476 conservatively modified variations. Every nucleic acid sequence herein which encodes a

polypeptide also describes silent variations of the nucleic acid. In certain contexts each codon in a nucleic acid (except AUG, which is ordinarily the only codon for methionine, and TGG, which is ordinarily the only codon for tryptophan) can be modified to yield a functionally similar molecule. Accordingly, a silent variation of a nucleic acid which encodes a polypeptide is implicit in a described sequence with respect to the expression product, but not necessarily with respect to actual probe sequences.

As to amino acid sequences, one of skill will recognize that individual substitutions, deletions or additions to a nucleic acid, peptide, polypeptide, or protein sequence which alters, adds or deletes a single amino acid or a small percentage of amino acids in the encoded sequence is a "conservatively modified variant" where the alteration results in the substitution of an amino acid with a chemically similar amino acid. Conservative substitution tables providing functionally similar amino acids are well known in the art. Such conservatively modified variants are in addition to and do not exclude polymorphic variants, interspecies homologs, and alleles of the invention. Typically conservative substitutions include for one another: 1) Alanine (A), Glycine (G); 2) Aspartic acid (D), Glutamic acid (E); 3) Asparagine (N), Glutamine (Q); 4) Arginine (R), Lysine (K); 5) Isoleucine (I), Leucine (L), Methionine (M), Valine (V); 6) Phenylalanine (F), Tyrosine (Y), Tryptophan (W); 7) Serine (S), Threonine (T); and 8) Cysteine (C), Methionine (M) (see, e.g., Creighton, Proteins (1984)).

Macromolecular structures such as polypeptide structures can be described in terms of various levels of organization. For a general discussion of this organization, see, e.g., Alberts, et al. (1994) Molecular Biology of the Cell (3rd ed.) and Cantor and Schimmel (1980) Biophysical Chemistry Part I: The Conformation of Biological Macromolecules. "Primary structure" refers to the amino acid sequence of a particular peptide. "Secondary structure" refers to locally ordered, three dimensional structures within a polypeptide. These structures are commonly known as domains. Domains are portions of a polypeptide that often form a compact unit of the polypeptide and are typically 25 to approximately 500 amino acids long. Typical domains are made up of sections of lesser organization such as stretches of β -sheet and α -helices. "Tertiary structure" refers to the complete three dimensional structure of a polypeptide monomer. "Quaternary structure" refers to the three dimensional structure formed, usually by the noncovalent association of independent tertiary units. Anisotropic terms are also known as energy terms.

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"Nucleic acid" or "oligonucleotide" or "polynucleotide" or grammatical equivalents used herein means at least two nucleotides covalently linked together. Oligonucleotides are typically from about 5, 6, 7, 8, 9, 10, 12, 15, 25, 30, 40, 50 or more nucleotides in length, up to about 100 nucleotides in length. Nucleic acids and polynucleotides are a polymers of any length, including longer lengths, e.g., 200, 300, 500, 1000, 2000, 3000, 5000, 7000, 10,000, etc. A nucleic acid of the present invention will generally contain phosphodiester bonds, although in some cases, nucleic acid analogs are included that may have at least one different linkage, e.g., phosphoramidate, phosphorothioate, phosphorodithioate, or Omethylphophoroamidite linkages (see Eckstein (1992) Oligonucleotides and Analogues: A Practical Approach Oxford University Press); and peptide nucleic acid backbones and linkages. Other analog nucleic acids include those with positive backbones; non-ionic backbones, and non-ribose backbones, including those described in U.S. Patent Nos. 5,235,033 and 5,034,506, and Chapters 6 and 7, in Sanghui and Cook, eds. Carbohydrate Modifications in Antisense Research, ASC Symposium Series 580. Nucleic acids containing one or more carbocyclic sugars are also included within one definition of nucleic acids. Modifications of the ribose-phosphate backbone may be done for a variety of reasons, e.g., to increase the stability and half-life of such molecules in physiological environments or as probes on a biochip. Mixtures of naturally occurring nucleic acids and analogs can be made; alternatively, mixtures of different nucleic acid analogs, and mixtures of naturally occurring nucleic acids and analogs may be made.

Particularly preferred are peptide nucleic acids (PNA) which includes peptide nucleic acid analogs. These backbones are substantially non-ionic under neutral conditions, in contrast to the highly charged phosphodiester backbone of naturally occurring nucleic acids. This results in two advantages. First, the PNA backbone exhibits improved hybridization kinetics. PNAs have larger changes in the melting temperature (T_m) for mismatched versus perfectly matched basepairs. DNA and RNA typically exhibit a 2-4° C drop in T_m for an internal mismatch. With the non-ionic PNA backbone, the drop is closer to 7-9° C. Similarly, due to their non-ionic nature, hybridization of the bases attached to these backbones is relatively insensitive to salt concentration. In addition, PNAs are not degraded by cellular enzymes, and thus can be more stable.

The nucleic acids may be single stranded or double stranded, as specified, or contain portions of both double stranded or single stranded sequence. As will be appreciated by those in the art, the depiction of a single strand also defines the sequence of the complementary

strand; thus the sequences described herein also provide the complement of the sequence. The nucleic acid may be DNA, both genomic and cDNA, RNA, or a hybrid, where the nucleic acid may contain combinations of deoxyribo- and ribo-nucleotides, and combinations of bases, including uracil, adenine, thymine, cytosine, guanine, inosine, xanthine hypoxanthine, isocytosine, isoguanine, etc. "Transcript" typically refers to a naturally occurring RNA, e.g., a pre-mRNA, hnRNA, or mRNA. As used herein, the term "nucleoside" includes nucleotides and nucleoside and nucleotide analogs, and modified nucleosides such as amino modified nucleosides. In addition, "nucleoside" includes non-naturally occurring analog structures. Thus, e.g., the individual units of a peptide nucleic acid, each containing a base, are referred to herein as a nucleoside.

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A "label" or a "detectable moiety" is a composition detectable by spectroscopic, photochemical, biochemical, immunochemical, physiological, chemical, or other physical means. For example, useful labels include ³²P, fluorescent dyes, electron-dense reagents, enzymes (e.g., as commonly used in an ELISA), biotin, digoxigenin, or haptens and proteins or other entities which can be made detectable, e.g., by incorporating a radiolabel into the peptide or used to detect antibodies specifically reactive with the peptide. The labels may be incorporated into the cancer nucleic acids, proteins, and antibodies. Many methods known in the art for conjugating the antibody to the label may be employed, including those methods described by Hunter, et al. (1962) Nature 144:945; David, et al. (1974) Biochemistry 13:1014-1021; Pain, et al. (1981) J. Immunol. Meth., 40:219-230; and Nygren (1982) J. Histochem. and Cytochem. 30:407-412.

An "effector" or "effector moiety" or "effector component" is a molecule that is bound (or linked, or conjugated), either covalently, through a linker or a chemical bond, or noncovalently, through ionic, van der Waals, electrostatic, or hydrogen bonds, to an antibody. The "effector" can be a variety of molecules including, e.g., detection moieties including radioactive compounds, fluorescent compounds, an enzyme or substrate, tags such as epitope tags, a toxin; activatable moieties, a chemotherapeutic agent; a lipase; an antibiotic; or a radioisotope emitting "hard" e.g., beta radiation.

A "labeled nucleic acid probe or oligonucleotide" is one that is bound, either covalently, through a linker or a chemical bond, or noncovalently, through ionic, van der Waals, electrostatic, or hydrogen bonds to a label such that the presence of the probe may be detected by detecting the presence of the label bound to the probe. Alternatively, method

WO 02/086443 PCT/US02/12476 using high affinity interactions may achieve the same results where one of a pair of binding

partners binds to the other, e.g., biotin, streptavidin.

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As used herein a "nucleic acid probe or oligonucleotide" is a nucleic acid capable of binding to a target nucleic acid of complementary sequence through one or more types of chemical bonds, usually through complementary base pairing, e.g., through hydrogen bond formation. As used herein, a probe may include natural (i.e., A, G, C, or T) or modified bases (7-deazaguanosine, inosine, etc.). In addition, the bases in a probe may be joined by a linkage other than a phosphodiester bond, preferably one that does not functionally interfere with hybridization. Thus, e.g., probes may be peptide nucleic acids in which the constituent bases are joined by peptide bonds rather than phosphodiester linkages. Probes may bind target sequences lacking complete complementarity with the probe sequence depending upon the stringency of the hybridization conditions. The probes are preferably directly labeled, e.g., with isotopes, chromophores, lumiphores, chromogens, or indirectly labeled, e.g., with biotin to which a streptavidin complex may later bind. By assaying for the presence or absence of the probe, one can detect the presence or absence of the select sequence or subsequence. Diagnosis or prognosis may be based at the genomic level, or at the level of RNA or protein expression.

The term "recombinant" when used with reference, e.g., to a cell, or nucleic acid, protein, or vector, indicates that the cell, nucleic acid, protein or vector, has been modified by the introduction of a heterologous nucleic acid or protein or the alteration of a native nucleic acid or protein, or that the cell is derived from a cell so modified. Thus, e.g., recombinant cells express genes that are not found within the native (non-recombinant) form of the cell or express native genes that are otherwise abnormally expressed, under expressed or not expressed at all. By the term "recombinant nucleic acid" herein is meant nucleic acid, originally formed in vitro, in general, by the manipulation of nucleic acid, e.g., using polymerases and endonucleases, in a form not normally found in nature. In this manner, operably linkage of different sequences is achieved. Thus an isolated nucleic acid, in a linear form, or an expression vector formed in vitro by ligating DNA molecules that are not normally joined, are both considered recombinant for the purposes of this invention. It is understood that once a recombinant nucleic acid is made and reintroduced into a host cell or organism, it will replicate non-recombinantly, i.e., using the in vivo cellular machinery of the host cell rather than in vitro manipulations; however, such nucleic acids, once produced recombinantly, although subsequently replicated non-recombinantly, are still considered

recombinant for the purposes of the invention. Similarly, a "recombinant protein" is a protein made using recombinant techniques, i.e., through the expression of a recombinant nucleic acid as depicted above.

The term "heterologous" when used with reference to portions of a nucleic acid indicates that the nucleic acid comprises two or more subsequences that are not normally found in the same relationship to each other in nature. For instance, the nucleic acid is typically recombinantly produced, having two or more sequences, e.g., from unrelated genes arranged to make a new functional nucleic acid, e.g., a promoter from one source and a coding region from another source. Similarly, a heterologous protein will often refer to two or more subsequences that are not found in the same relationship to each other in nature (e.g., a fusion protein).

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A "promoter" is typically an array of nucleic acid control sequences that direct transcription of a nucleic acid. As used herein, a promoter includes necessary nucleic acid sequences near the start site of transcription, such as, in the case of a polymerase II type promoter, a TATA element. A promoter also optionally includes distal enhancer or repressor elements, which can be located as much as several thousand base pairs from the start site of transcription. A "constitutive" promoter is a promoter that is active under most environmental and developmental conditions. An "inducible" promoter is a promoter that is active under environmental or developmental regulation. The term "operably linked" refers to a functional linkage between a nucleic acid expression control sequence (such as a promoter, or array of transcription factor binding sites) and a second nucleic acid sequence, e.g., wherein the expression control sequence directs transcription of the nucleic acid corresponding to the second sequence.

An "expression vector" is a nucleic acid construct, generated recombinantly or synthetically, with a series of specified nucleic acid elements that permit transcription of a particular nucleic acid in a host cell. The expression vector can be part of a plasmid, virus, or nucleic acid fragment. Typically, the expression vector includes a nucleic acid to be transcribed in operable linkage to a promoter.

The phrase "selectively (or specifically) hybridizes to" refers to the binding, duplexing, or hybridizing of a molecule selectively to a particular nucleotide sequence under stringent hybridization conditions when that sequence is present in a complex mixture (e.g., total cellular or library DNA or RNA).

The phrase "stringent hybridization conditions" refers to conditions under which a probe will hybridize to its target subsequence, typically in a complex mixture of nucleic acids, but to essentially no other sequences. Stringent conditions are sequence-dependent and will be different in different circumstances. Longer sequences hybridize specifically at higher temperatures. An extensive guide to the hybridization of nucleic acids is found in . 5 "Overview of principles of hybridization and the strategy of nucleic acid assays" in Tijssen (1993) Techniques in Biochemistry and Molecular Biology--Hybridization with Nucleic Probes (vol. 24) Elsevier. Generally, stringent conditions are selected to be about 5-10° C lower than the thermal melting point (T_m) for the specific sequence at a defined ionic strength pH. The T_m is the temperature (under defined ionic strength, pH, and nucleic concentration) 10 at which 50% of the probes complementary to the target hybridize to the target sequence at equilibrium (as the target sequences are present in excess, at T_m, 50% of the probes are occupied at equilibrium). Stringent conditions will be those in which the salt concentration is less than about 1.0 M sodium ion, typically about 0.01 to 1.0 M sodium ion concentration (or other salts) at pH 7.0 to 8.3 and the temperature is at least about 30° C for short probes (e.g., 15 10 to 50 nucleotides) and at least about 60° C for long probes (e.g., greater than 50 nucleotides). Stringent conditions may also be achieved with the addition of destabilizing agents such as formamide. For selective or specific hybridization, a positive signal is typically at least two times background, preferably 10 times background hybridization. Exemplary stringent hybridization conditions are often: 50% formamide, 5x SSC, and 1% 20 SDS, incubating at 42° C, or, 5x SSC, 1% SDS, incubating at 65° C, with wash in 0.2x SSC, and 0.1% SDS at 65° C. For PCR, a temperature of about 36° C is typical for low stringency amplification, although annealing temperatures may vary between about 32° C and 48° C depending on primer length. For high stringency PCR amplification, a temperature of about 62° C is typical, although high stringency annealing temperatures can range from about 50° C 25 to about 65° C, depending on the primer length and specificity. Typical cycle conditions for both high and low stringency amplifications include a denaturation phase of 90° C - 95° C for 0.5 - 2 min., an annealing phase lasting 0.5 - 2 min., and an extension phase of about 72° C for 1 - 2 min. Protocols and guidelines for low and high stringency amplification reactions are provided, e.g., in Innis, et al.(1990) PCR Protocols, A Guide to Methods and 30 Applications.

Nucleic acids that do not hybridize to each other under stringent conditions are still substantially identical if the polypeptides which they encode are substantially identical. This

WO 02/086443 PCT/US02/12476 occurs, e.g., when a copy of a nucleic acid is created using the maximum codon degeneracy

permitted by the genetic code. In such cases, the nucleic acids typically hybridize under moderately stringent hybridization conditions. Exemplary "moderately stringent hybridization conditions" include a hybridization in a buffer of 40% formamide, 1 M NaCl, 1% SDS at 37° C, and a wash in 1X SSC at 45° C. A positive hybridization is at least twice background. Alternative hybridization and wash conditions can be utilized to provide conditions of similar stringency. Additional guidelines for determining hybridization parameters are provided in numerous reference, e.g., Ausubel, et al. (ed.) <u>Current Protocols in Molecular Biology</u> Lippincott.

The phrase "functional effects" in the context of assays for testing compounds that modulate activity of a lung cancer protein includes the determination of a parameter that is indirectly or directly under the influence of the lung cancer protein or nucleic acid, e.g., a physiological, enzymatic, functional, physical, or chemical effect, such as the ability to decrease lung cancer. It includes ligand binding activity; cell viability, cell growth on soft agar; anchorage dependence; contact inhibition and density limitation of growth; cellular proliferation; cellular transformation; growth factor or serum dependence; tumor specific marker levels; invasiveness into Matrigel; tumor growth and metastasis in vivo; mRNA and protein expression in cells undergoing metastasis, and other characteristics of lung cancer cells. "Functional effects" include in vitro, in vivo, and ex vivo activities.

By "determining the functional effect" is meant assaying for a compound that increases or decreases a parameter that is indirectly or directly under the influence of a lung cancer protein sequence, e.g., physiological, functional, enzymatic, physical, or chemical effects. Such functional effects can be measured by many means known to those skilled in the art, e.g., changes in spectroscopic characteristics (e.g., fluorescence, absorbance, refractive index), hydrodynamic (e.g., shape), chromatographic, or solubility properties for the protein, measuring inducible markers or transcriptional activation of the lung cancer protein; measuring binding activity or binding assays, e.g., binding to antibodies or other ligands, and measuring cellular proliferation. Determination of the functional effect of a compound on lung cancer can also be performed using lung cancer assays known to those of skill in the art such as an *in vitro* assays, e.g., cell growth on soft agar; anchorage dependence; contact inhibition and density limitation of growth; cellular proliferation; cellular transformation; growth factor or serum dependence; tumor specific marker levels; invasiveness into Matrigel; tumor growth and metastasis *in vivo*; mRNA and protein

expression in cells undergoing metastasis, and other characteristics of lung cancer cells. The functional effects can be evaluated by many means known to those skilled in the art, e.g., microscopy for quantitative or qualitative measures of alterations in morphological features, measurement of changes in RNA or protein levels for lung cancer-associated sequences, measurement of RNA stability, identification of downstream or reporter gene expression (CAT, luciferase, β -gal, GFP, and the like), e.g., via chemiluminescence, fluorescence, colorimetric reactions, antibody binding, inducible markers, and ligand binding assays.

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"Inhibitors". "activators", and "modulators" of lung cancer polynucleotide and polypeptide sequences are used to refer to activating, inhibitory, or modulating molecules or compounds identified using in vitro and in vivo assays of lung cancer polynucleotide and polypeptide sequences. Inhibitors are compounds that, e.g., bind to, partially or totally block activity, decrease, prevent, delay activation, inactivate, desensitize, or down regulate the activity or expression of lung cancer proteins, e.g., antagonists. Antisense or inhibitory nucleic acids may seem to inhibit expression and subsequent function of the protein. "Activators" are compounds that increase, open, activate, facilitate, enhance activation, sensitize, agonize, or up regulate lung cancer protein activity. Inhibitors, activators, or modulators also include genetically modified versions of lung cancer proteins, e.g., versions with altered activity, as well as naturally occurring and synthetic ligands, antagonists, agonists, antibodies, small chemical molecules and the like. Such assays for inhibitors and activators include, e.g., expressing the lung cancer protein in vitro, in cells, or cell membranes, applying putative modulator compounds, and then determining the functional effects on activity, as described above. Activators and inhibitors of lung cancer can also be identified by incubating lung cancer cells with the test compound and determining increases or decreases in the expression of 1 or more lung cancer proteins, e.g., 1, 2, 3, 4, 5, 10, 15, 20, 25, 30, 40, 50 or more lung cancer proteins, such as lung cancer proteins encoded by the sequences set out in Tables 1A-16.

Samples or assays comprising lung cancer proteins that are treated with a potential activator, inhibitor, or modulator are compared to control samples without the inhibitor, activator, or modulator to examine the extent of inhibition. Control samples (untreated with inhibitors) are assigned a relative protein activity value of 100%. Inhibition of a polypeptide is achieved when the activity value relative to the control is about 80%, preferably 50%, more preferably 25-0%. Activation of a lung cancer polypeptide is achieved when the activity value relative to the control (untreated with activators) is 110%, more preferably 150%, more

WO 02/086443 PCT/US02/12476 preferably 200-500% (i.e., two to five fold higher relative to the control), more preferably

1000-3000% higher.

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The phrase "changes in cell growth" refers to any change in cell growth and proliferation characteristics *in vitro* or *in vivo*, such as cell viability, formation of foci, anchorage independence, semi-solid or soft agar growth, changes in contact inhibition and density limitation of growth, loss of growth factor or serum requirements, changes in cell morphology, gaining or losing immortalization, gaining or losing tumor specific markers, ability to form or suppress tumors when injected into suitable animal hosts, and/or immortalization of the cell. See, e.g., Freshney (1994) <u>Culture of Animal Cells a Manual of Basic Technique</u> pp. 231-241 (3rd ed.).

"Tumor cell" refers to precancerous, cancerous, and normal cells in a tumor.

"Cancer cells," "transformed" cells, or "transformation" in tissue culture, refers to spontaneous or induced phenotypic changes that do not necessarily involve the uptake of new genetic material. Although transformation can arise from infection with a transforming virus and incorporation of new genomic DNA, or uptake of exogenous DNA, it can also arise spontaneously or following exposure to a carcinogen, thereby mutating an endogenous gene. Transformation is associated with phenotypic changes, such as immortalization of cells, aberrant growth control, nonmorphological changes, and/or malignancy (see, Freshney (1994) Culture of Animal Cells a Manual of Basic Technique (3rd ed.)).

"Antibody" refers to a polypeptide comprising a framework region from an immunoglobulin gene or fragments thereof that specifically binds and recognizes an antigen. The recognized immunoglobulin genes include the kappa, lambda, alpha, gamma, delta, epsilon, and mu constant region genes, as well as the myriad immunoglobulin variable region genes. Light chains are classified as either kappa or lambda. Heavy chains are classified as gamma, mu, alpha, delta, or epsilon, which in turn define the immunoglobulin classes, IgG, IgM, IgA, IgD, and IgE, respectively. Typically, the antigen-binding region of an antibody or its functional equivalent will be most critical in specificity and affinity of binding. See Paul, Fundamental Immunology.

An exemplary immunoglobulin (antibody) structural unit comprises a tetramer. Each tetramer is composed of two identical pairs of polypeptide chains, each pair having one "light" (about 25 kD) and one "heavy" chain (about 50-70 kD). The N-terminus of each chain defines a variable region of about 100 to 110 or more amino acids primarily responsible

WO 02/086443 PCT/US02/12476 for antigen recognition. The terms variable light chain (V_L) and variable heavy chain (V_H)

refer to these light and heavy chains respectively.

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Antibodies exist, e.g., as intact immunoglobulins or as a number of well-characterized fragments produced by digestion with various peptidases. Thus, e.g., pepsin digests an antibody below the disulfide linkages in the hinge region to produce F(ab)'₂, a dimer of Fab which itself is a light chain joined to V_H-C_H1 by a disulfide bond. The F(ab)'₂ may be reduced under mild conditions to break the disulfide linkage in the hinge region, thereby converting the F(ab)'₂ dimer into an Fab' monomer. The Fab' monomer is essentially Fab with part of the hinge region (see Paul (ed. 1999) Fundamental Immunology (4th ed.). While various antibody fragments are defined in terms of the digestion of an intact antibody, one of skill will appreciate that such fragments may be synthesized *de novo* either chemically or by using recombinant DNA methodology. Thus, the term antibody, as used herein, also includes antibody fragments either produced by the modification of whole antibodies, or those synthesized *de novo* using recombinant DNA methodologies (e.g., single chain Fv) or those identified using phage display libraries (see, e.g., McCafferty, et al. (1990) Nature 348:552-554).

For preparation of antibodies, e.g., recombinant, monoclonal, or polyclonal antibodies, many technique known in the art can be used (see, e.g., Kohler and Milstein (1975) Nature 256:495-497; Kozbor, et al. (1983) Immunology Today 4:72; Cole, et al. (1985), pp. 77-96 in Monoclonal Antibodies and Cancer Therapy; Coligan (1991 and supplements) Current Protocols in Immunology; Harlow and Lane (1988) Antibodies, A Laboratory Manual; and Goding (1986) Monoclonal Antibodies: Principles and Practice (2d ed.)). Techniques for the production of single chain antibodies (U.S. Patent 4,946,778) can be adapted to produce antibodies to polypeptides of this invention. Also, transgenic mice, or other organisms such as other mammals, may be used to express humanized antibodies. Alternatively, phage display technology can be used to identify antibodies and heteromeric Fab fragments that specifically bind to selected antigens (see, e.g., McCafferty, et al. (1990) Nature 348:552-554; Marks, et al. (1992) Biotechnology 10:779-783).

A "chimeric antibody" is an antibody molecule in which, e.g, (a) the constant region, or a portion thereof, is altered, replaced, or exchanged so that the antigen binding site (variable region) is linked to a constant region of a different or altered class, effector function, and/or species, or an entirely different molecule which confers new properties to the chimeric antibody, e.g., an enzyme, toxin, hormone, growth factor, drug, etc.; or (b) the

variable region, or a portion thereof, is altered, replaced, or exchanged with a variable region having a different or altered antigen specificity.

Identification of lung cancer-associated sequences

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In one aspect, the expression levels of genes are determined in different patient samples for which diagnosis information is desired, to provide expression profiles. An expression profile of a particular sample is essentially a "fingerprint" of the state of the sample; while two states may have any particular gene similarly expressed, the evaluation of a number of genes simultaneously allows the generation of a gene expression profile that is characteristic of the state of the cell. That is, normal tissue may be distinguished from cancerous or metastatic cancerous tissue, or metastatic cancerous tissue can be compared with tissue from surviving cancer patients. By comparing expression profiles of tissue in known different lung cancer states, information regarding which genes are important (including both up- and down-regulation of genes) in each of these states is obtained. Molecular profiling may distinguish subtypes of a currently collective disease designation, e.g., different forms of lung cancer (chronic disease, adenocarcinoma, etc.)

The identification of sequences that are differentially expressed in lung cancer versus non-lung cancer tissue allows the use of this information in a number of ways. For example, a particular treatment regime may be evaluated: does a chemotherapeutic drug act to downregulate lung cancer, and thus tumor growth or recurrence, in a particular patient. Alternatively, a treatment step may induce other markers which may be used as targets to destroy tumor cells. Similarly, diagnosis and treatment outcomes may be done or confirmed by comparing patient samples with the known expression profiles. Malignant diseasemay be compared to non-malignant conditions. Metastatic tissue can also be analyzed to determine the stage of lung cancer in the tissue, or origin of primary tumor, e.g., metastasis from a remote primary site. Furthermore, these gene expression profiles (or individual genes) allow screening of drug candidates with an eye to mimicking or altering a particular expression profile; e.g., screening can be done for drugs that suppress the lung cancer expression profile. This may be done by making biochips comprising sets of the important lung cancer genes, which can then be used in these screens. PCR methods may be applied with selected primer pairs, and analysis may be of RNA or of genomic sequences. These methods can also be done on the protein basis; that is, protein expression levels of the lung cancer proteins can be evaluated for diagnostic purposes or to screen candidate agents. In addition, the lung cancer

nucleic acid sequences can be administered for gene therapy purposes, including the administration of antisense nucleic acids, or the lung cancer proteins (including antibodies and other modulators thereof) administered as therapeutic drugs or as protein or DNA vaccines.

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Thus the present invention provides nucleic acid and protein sequences that are differentially expressed in lung cancer relative to normal tissues and/or non-malignant lung disease, or in different types of lung disease, herein termed "lung cancer sequences." As outlined below, lung cancer sequences include those that are up-regulated (i.e., expressed at a higher level) in lung cancer, as well as those that are down-regulated (i.e., expressed at a lower level). In a preferred embodiment, the lung cancer sequences are from humans; however, as will be appreciated by those in the art, lung cancer sequences from other organisms may be useful in animal models of disease and drug evaluation; thus, other lung cancer sequences are provided, from vertebrates, including mammals, including rodents (rats, mice, hamsters, guinea pigs, etc.), primates, farm animals (including sheep, goats, pigs, cows, horses, etc.) and pets (dogs, cats, etc.). Lung cancer sequences from other organisms may be obtained using the techniques outlined below.

Lung cancer sequences can include both nucleic acid and amino acid sequences. As will be appreciated by those in the art and is more fully outlined below, lung cancer nucleic acid sequences are useful in a variety of applications, including diagnostic applications, which will detect naturally occurring nucleic acids, as well as screening applications; e.g., biochips comprising nucleic acid probes or PCR microtiter plates with selected probes to the lung cancer sequences can be generated.

A lung cancer sequence can be initially identified by substantial nucleic acid and/or amino acid sequence homology to the lung cancer sequences outlined herein. Such homology can be based upon the overall nucleic acid or amino acid sequence, and is generally determined as outlined below, e.g., using homology programs or hybridization conditions.

For identifying lung cancer-associated sequences, the lung cancer screen typically includes comparing genes identified in different tissues, e.g., normal and cancerous tissues, cancer and non-malignant conditions, non-malignant conditions and normal tissues, or tumor tissue samples from patients who have metastatic disease vs. non metastatic tissue. Other suitable tissue comparisons include comparing lung cancer samples with metastatic cancer samples from other cancers, such as, breast, other gastrointestinal cancers, prostate, ovarian,

etc. Samples of, non metastatic disease tissue and tissue undergoing metastasis are applied to biochips comprising nucleic acid probes. The samples are first microdissected, if applicable, and treated as is known in the art for the preparation of mRNA. Suitable biochips are commercially available, e.g., from Affymetrix, Santa Clara, CA. Gene expression profiles as described herein are generated and the data analyzed.

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In one embodiment, the genes showing changes in expression as between normal and disease states are compared to genes expressed in other normal tissues, preferably normal lung, but also including, and not limited to colon, heart, brain, liver, breast, kidney, muscle, prostate, small intestine, large intestine, spleen, bone, and/or placenta. In a preferred embodiment, those genes identified during the lung cancer screen that are expressed in significant amounts in other tissues (e.g., essential organs) are removed from the profile, although in some embodiments, this is not necessary (e.g., where organs may be dispensible at a later stage of life). That is, when screening for drugs, it is usually preferable that the target expression be disease specific, to minimize possible side effects on other organs.

In a preferred embodiment, lung cancer sequences are those that are up-regulated in lung cancer; that is, the expression of these genes is higher in cancerous tissue than in normal lung or other tissue. "Up-regulation" as used herein means, when the ratio is presented as a number greater than one, that the ratio is greater than one, preferably 1.5 or greater, more preferably 2.0 or greater. Another embodiment is directed to sequences up-regulated in nonmalignant conditions relative to normal. Unigene cluster identification numbers and accession numbers herein are for the GenBank sequence database and the sequences of the accession numbers are hereby expressly incorporated by reference. GenBank is known in the art, see, e.g., Benson, DA, et al (1998) Nucleic Acids Research 26:1-7 and http://www.ncbi.nlm.nih.gov/. Sequences are also available in other databases, e.g., European Molecular Biology Laboratory (EMBL) and DNA Database of Japan (DDBJ). Another embodiment is directed to sequences up-regulated in non-malignant conditions relative to normal. In some situations, the sequences may be derived from assembly of available sequences or be predicted from genomic DNA using exon prediction algorithms, such as FGENESH (Salamov and Solovyev (2000) Genome Res. 10:516-522). In other situations, sequences have been derived from cloning and sequencing of isolated nucleic acids.

In another preferred embodiment, lung cancer sequences are those that are downregulated in the lung cancer; that is, the expression of these genes is lower in cancerous tissue

or normal lung or other tissue. "Down-regulation" as used herein means, when the ratio is presented as a number greater than one, that the ratio is greater than one, preferably 1.5 or greater, more preferably 2.0 or greater, or, when the ratio is presented as a number less than one, that the ratio is less than one, preferably 0.5 or less, more preferably 0.25 or less.

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Informatics

The ability to identify genes that are over or under expressed in lung cancer can additionally provide high-resolution, high-sensitivity datasets which can be used in the areas of diagnostics, therapeutics, drug development, pharmacogenetics, protein structure, biosensor development, and other related areas. For example, the expression profiles can be used in diagnostic or prognostic evaluation of patients with lung cancer. Or as another example, subcellular toxicological information can be generated to better direct drug structure and activity correlation (see Anderson (1998) Pharmaceutical Proteomics: Targets,

Mechanism, and Function, paper presented at the IBC Proteomics conference, Coronado, CA (June 11-12, 1998)). Subcellular toxicological information can also be utilized in a biological sensor device to predict the likely toxicological effect of chemical exposures and likely tolerable exposure thresholds (see U.S. Patent No. 5,811,231). Similar advantages accrue from datasets relevant to other biomolecules and bioactive agents (e.g., nucleic acids, saccharides, lipids, drugs, and the like).

Thus, in another embodiment, the present invention provides a database that includes at least one set of assay data. The data contained in the database is acquired, e.g., using array analysis either singly or in a library format. The database can be in a form in which data can be maintained and transmitted, but is preferably an electronic database. The electronic database of the invention can be maintained on any electronic device allowing for the storage of and access to the database, such as a personal computer, but is preferably distributed on a wide area network, such as the World Wide Web.

The focus of the present section on databases that include peptide sequence data is for clarity of illustration only. It will be apparent to those of skill in the art that similar databases can be assembled for assay data acquired using an assay of the invention.

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The compositions and methods for identifying and/or quantitating the relative and/or absolute abundance of a variety of molecular and macromolecular species from a biological sample representing lung cancer, i.e., the identification of lung cancer-associated sequences described herein, provide an abundance of information, which can be correlated with

WO 02/086443 PCT/US02/12476 pathological conditions, predisposition to disease, drug testing, therapeutic monitoring, gene-

disease causal linkages, identification of correlates of immunity and physiological status, among others. Although the data generated from the assays of the invention is suited for manual review and analysis, in a preferred embodiment, data processing using high-speed

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An array of methods for indexing and retrieving biomolecular information is known in the art. For example, U.S. Patents 6,023,659 and 5,966,712 disclose a relational database system for storing biomolecular sequence information in a manner that allows sequences to be catalogued and searched according to one or more protein function hierarchies. U.S. Patent 5,953,727 discloses a relational database having sequence records containing information in a format that allows a collection of partial-length DNA sequences to be catalogued and searched according to association with one or more sequencing projects for obtaining full-length sequences from the collection of partial length sequences. U.S. Patent 5,706,498 discloses a gene database retrieval system for making a retrieval of a gene sequence similar to a sequence data item in a gene database based on the degree of similarity between a key sequence and a target sequence. U.S. Patent 5,538,897 discloses a method using mass spectroscopy fragmentation patterns of peptides to identify amino acid sequences in computer databases by comparison of predicted mass spectra with experimentally-derived mass spectra using a closeness-of-fit measure. U.S. Patent 5,926,818 discloses a multidimensional database comprising a functionality for multi-dimensional data analysis described as on-line analytical processing (OLAP), which entails the consolidation of projected and actual data according to more than one consolidation path or dimension. U.S. Patent 5,295,261 reports a hybrid database structure in which the fields of each database record are divided into two classes, navigational and informational data, with navigational fields stored in a hierarchical topological map which can be viewed as a tree structure or as the merger of two or more such tree structures.

See also Mount, et al. (2001) Bioinformatics; Durbin, et al. (eds., 1999) Biological Sequence Analysis: Probabilistic Models of Proteins and Nucleic Acids (; Baxevanis and Oeullette (eds., 1998) Bioinformatics: A Practical Guide to the Analysis of Genes and Proteins); Rashidi and Buehler (1999) Bioinformatics: Basic Applications in Biological Science and Medicine; Setubal, et al. (eds 1997) Introduction to Computational Molecular Biology; Misener and Krawetz (eds, 2000) Bioinformatics: Methods and Protocols; Higgins and Taylor (eds., 2000) Bioinformatics: Sequence, Structure, and Databanks: A Practical

WO 02/086443 PCT/US02/12476 Approach; Brown (2001) Bioinformatics: A Biologist's Guide to Biocomputing and the

Internet; Han and Kamber (2000) Data Mining: Concepts and Techniques (2000); and Waterman (1995) Introduction to Computational Biology: Maps, Sequences, and Genomes.

The present invention provides a computer database comprising a computer and software for storing in computer-retrievable form assay data records cross-tabulated, e.g., with data specifying the source of the target-containing sample from which each sequence specificity record was obtained.

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In an exemplary embodiment, at least one of the sources of target-containing sample is from a control tissue sample known to be free of pathological disorders. In a variation, at least one of the sources is a known pathological tissue specimen, e.g., a neoplastic lesion or another tissue specimen to be analyzed for lung cancer. In another variation, the assay records cross-tabulate one or more of the following parameters for each target species in a sample: (1) a unique identification code, which can include, e.g., a target molecular structure and/or characteristic separation coordinate (e.g., electrophoretic coordinates); (2) sample source; and (3) absolute and/or relative quantity of the target species present in the sample.

The invention also provides for the storage and retrieval of a collection of target data in a computer data storage apparatus, which can include magnetic disks, optical disks, magneto-optical disks, DRAM, SRAM, SGRAM, SDRAM, RDRAM, DDR RAM, magnetic bubble memory devices, and other data storage devices, including CPU registers and on-CPU data storage arrays. Typically, the target data records are stored as a bit pattern in an array of magnetic domains on a magnetizable medium or as an array of charge states or transistor gate states, such as an array of cells in a DRAM device (e.g., each cell comprised of a transistor and a charge storage area, which may be on the transistor). In one embodiment, the invention provides such storage devices, and computer systems built therewith, comprising a bit pattern encoding a protein expression fingerprint record comprising unique identifiers for at least 10 target data records cross-tabulated with target source.

When the target is a peptide or nucleic acid, the invention preferably provides a method for identifying related peptide or nucleic acid sequences, comprising performing a computerized comparison between a peptide or nucleic acid sequence assay record stored in or retrieved from a computer storage device or database and at least one other sequence. The comparison can include a sequence analysis or comparison algorithm or computer program embodiment thereof (e.g., FASTA, TFASTA, GAP, BESTFIT) and/or the comparison may

be of the relative amount of a peptide or nucleic acid sequence in a pool of sequences determined from a polypeptide or nucleic acid sample of a specimen.

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The invention also preferably provides a magnetic disk, such as an IBM-compatible (DOS, Windows, Windows95/98/2000, Windows NT, OS/2) or other format (e.g., Linux, SunOS, Solaris, AIX, SCO Unix, VMS, MV, Macintosh, etc.) floppy diskette or hard (fixed, Winchester) disk drive, comprising a bit pattern encoding data from an assay of the invention in a file format suitable for retrieval and processing in a computerized sequence analysis, comparison, or relative quantitation method.

The invention also provides a network, comprising a plurality of computing devices linked via a data link, such as an Ethernet cable (coax or 10BaseT), telephone line, ISDN line, wireless network, optical fiber, or other suitable signal transmission medium, whereby at least one network device (e.g., computer, disk array, etc.) comprises a pattern of magnetic domains (e.g., magnetic disk) and/or charge domains (e.g., an array of DRAM cells) composing a bit pattern encoding data acquired from an assay of the invention.

The invention also provides a method for transmitting assay data that includes generating an electronic signal on an electronic communications device, such as a modem, ISDN terminal adapter, DSL, cable modem, ATM switch, or the like, wherein the signal includes (in native or encrypted format) a bit pattern encoding data from an assay or a database comprising a plurality of assay results obtained by the method of the invention.

In a preferred embodiment, the invention provides a computer system for comparing a query target to a database containing an array of data structures, such as an assay result obtained by the method of the invention, and ranking database targets based on the degree of identity and gap weight to the target data. A central processor is preferably initialized to load and execute the computer program for alignment and/or comparison of the assay results. Data for a query target is entered into the central processor via an I/O device. Execution of the computer program results in the central processor retrieving the assay data from the data file, which comprises a binary description of an assay result.

The target data or record and the computer program can be transferred to secondary memory, which is typically random access memory (e.g., DRAM, SRAM, SGRAM, or SDRAM). Targets are ranked according to the degree of correspondence between a selected assay characteristic (e.g., binding to a selected affinity moiety) and the same characteristic of the query target and results are output via an I/O device. For example, a central processor can be a conventional computer (e.g., Intel Pentium, PowerPC, Alpha, PA-8000, SPARC,

MIPS 4400, MIPS 10000, VAX, etc.); a program can be a commercial or public domain molecular biology software package (e.g., UWGCG Sequence Analysis Software, Darwin); a data file can be an optical or magnetic disk, a data server, a memory device (e.g., DRAM, SRAM, SGRAM, SDRAM, EPROM, bubble memory, flash memory, etc.); an I/O device can be a terminal comprising a video display and a keyboard, a modem, an ISDN terminal adapter, an Ethernet port, a punched card reader, a magnetic strip reader, or other suitable I/O device.

The invention also preferably provides the use of a computer system, such as that described above, which comprises: (1) a computer; (2) a stored bit pattern encoding a collection of peptide sequence specificity records obtained by the methods of the invention, which may be stored in the computer; (3) a comparison target, such as a query target; and (4) a program for alignment and comparison, typically with rank-ordering of comparison results on the basis of computed similarity values.

Characteristics of lung cancer-associated proteins

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Lung cancer proteins of the present invention may be classified as secreted proteins, transmembrane proteins or intracellular proteins. In one embodiment, the lung cancer protein is an intracellular protein. Intracellular proteins may be found in the cytoplasm and/or in the nucleus. Intracellular proteins are involved in all aspects of cellular function and replication (including, e.g., signaling pathways); aberrant expression of such proteins often results in unregulated or disregulated cellular processes (see, e.g., Alberts (ed. 1994) Molecular Biology of the Cell (3d ed.). For example, many intracellular proteins have enzymatic activity such as protein kinase activity, protein phosphatase activity, protease activity, nucleotide cyclase activity, polymerase activity and the like. Intracellular proteins also serve as docking proteins that are involved in organizing complexes of proteins, or targeting proteins to various subcellular localizations, and are involved in maintaining the structural integrity of organelles.

An increasingly appreciated concept in characterizing proteins is the presence in the proteins of one or more structural motifs for which defined functions have been attributed. In addition to the highly conserved sequences found in the enzymatic domain of proteins, highly conserved sequences have been identified in proteins that are involved in protein-protein interaction. For example, Src-homology-2 (SH2) domains bind tyrosine-phosphorylated targets in a sequence dependent manner. PTB domains, which are distinct from SH2

domains, also bind tyrosine phosphorylated targets. SH3 domains bind to proline-rich targets. In addition, PH domains, tetratricopeptide repeats and WD domains to name only a few, have been shown to mediate protein-protein interactions. Some of these may also be involved in binding to phospholipids or other second messengers. As will be appreciated by one of ordinary skill in the art, these motifs can be identified on the basis of amino acid sequence; thus, an analysis of the sequence of proteins may provide insight into both the enzymatic potential of the molecule and/or molecules with which the protein may associate. One useful database is Pfam (protein families), which is a large collection of multiple sequence alignments and hidden Markov models covering many common protein domains. Versions are available via the internet from Washington University in St. Louis, the Sanger Center in England, and the Karolinska Institute in Sweden (see, e.g., Bateman, et al (2000) Nuc. Acids Res. 28:263-266; Sonnhammer, et al. (1997) Proteins 28:405-420; Bateman, et al. (1999) Nuc. Acids Res. 27:260-262; and Sonnhammer, et al. (1998) Nuc. Acids Res. 26:320-322).

In another embodiment, the lung cancer sequences are transmembrane proteins. Transmembrane proteins are molecules that span a phospholipid bilayer of a cell. They may have an intracellular domain, an extracellular domain, or both. The intracellular domains of such proteins may have a number of functions including those already described for intracellular proteins. For example, the intracellular domain may have enzymatic activity and/or may serve as a binding site for additional proteins. Frequently the intracellular domain of transmembrane proteins serves both roles. For example certain receptor tyrosine kinases have both protein kinase activity and SH2 domains. In addition, autophosphorylation of tyrosines on the receptor molecule itself, creates binding sites for additional SH2 domain containing proteins.

Transmembrane proteins may contain from one to many transmembrane domains. For example, receptor tyrosine kinases, certain cytokine receptors, receptor guanylyl cyclases and receptor serine/threonine protein kinases contain a single transmembrane domain. However, various other proteins including channels, pumps, and adenylyl cyclases contain numerous transmembrane domains. Many important cell surface receptors such as G protein coupled receptors (GPCRs) are classified as "seven transmembrane domain" proteins, as they contain 7 membrane spanning regions. Characteristics of transmembrane domains include approximately 17 consecutive hydrophobic amino acids that may be followed by charged amino acids. Therefore, upon analysis of the amino acid sequence of a particular protein, the

localization and number of transmembrane domains within the protein may be predicted (see, e.g., PSORT web site http://psort.nibb.ac.jp/).

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The extracellular domains of transmembrane proteins are diverse; however, conserved motifs are found repeatedly among various extracellular domains. Conserved structure and/or functions have been ascribed to different extracellular motifs. Many extracellular domains are involved in binding to other molecules. In one aspect, extracellular domains are found on receptors. Factors that bind the receptor domain include circulating ligands, which may be peptides, proteins, or small molecules such as adenosine and the like. For example, growth factors such as EGF, FGF, and PDGF are circulating growth factors that bind to their cognate receptors to initiate a variety of cellular responses. Other factors include cytokines, mitogenic factors, hormones, neurotrophic factors and the like. Extracellular domains also bind to cell-associated molecules. In this respect, they may mediate cell-cell interactions. Cell-associated ligands can be tethered to the cell, e.g., via a glycosylphosphatidylinositol (GPI) anchor, or may themselves be transmembrane proteins. Extracellular domains may also associate with the extracellular matrix and contribute to the maintenance of the cell structure.

Lung cancer proteins that are transmembrane are particularly preferred in the present invention as they are readily accessible targets for extracellular immunotherapeutics, as are described herein. In addition, as outlined below, transmembrane proteins can be also useful in imaging modalities. Antibodies may be used to label such readily accessible proteins in situ or in histological analysis. Alternatively, antibodies can also label intracellular proteins, in which case analytical samples are typically permeablized to provide access to intracellular proteins. In addition, some membrane proteins can be processed to release a soluble protein, or to expose a residual fragment. Released soluble proteins may be useful diagnostic markers, processed residual protein fragments may be useful lung markers of disease.

It will also be appreciated by those in the art that a transmembrane protein can be made soluble by removing transmembrane sequences, e.g., through recombinant methods. Furthermore, transmembrane proteins that have been made soluble can be made to be secreted through recombinant means by adding an appropriate signal sequence.

In another embodiment, the lung cancer proteins are secreted proteins; the secretion of which can be either constitutive or regulated. These proteins may have a signal peptide or signal sequence that targets the molecule to the secretory pathway. Secreted proteins are involved in numerous physiological events; e.g., if circulating, they often serve to transmit

signals to various other cell types. The secreted protein may function in an autocrine manner (acting on the cell that secreted the factor), a paracrine manner (acting on cells in close proximity to the cell that secreted the factor), an endocrine manner (acting on cells at a distance, e.g., secretion into the blood stream), or exocrine (secretion, e.g., through a duct or to adjacent epithelial surface as sweat glands, sebaceous glands, pancreatic ducts, lacrimal glands, mammary glands, sax producing glands of the ear, etc.). Thus secreted molecules often find use in modulating or altering numerous aspects of physiology. Lung cancer proteins that are secreted proteins are particularly preferred in the present invention as they serve as good targets for diagnostic markers, e.g., for blood, plasma, serum, or stool tests. Those which are enzymes may be antibody or small molecule targets. Others may be useful as vaccine targets, e.g., via CTL mechanisms.

Use of lung cancer nucleic acids

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As described above, lung cancer sequence is initially identified by substantial nucleic acid and/or amino acid sequence homology or linkage to the lung cancer sequences outlined herein. Such homology can be based upon the overall nucleic acid or amino acid sequence, and is generally determined as outlined below, using either homology programs or hybridization conditions. Typically, linked sequences on a mRNA are found on the same molecule.

The lung cancer nucleic acid sequences of the invention, e.g., the sequences in Tables 1A-16, can be fragments of larger genes, i.e., they are nucleic acid segments. "Genes" in this context includes coding regions, non-coding regions, and mixtures of coding and non-coding regions. Accordingly, as will be appreciated by those in the art, using the sequences provided herein, extended sequences, in either direction, of the lung cancer genes can be obtained, using techniques well known in the art for cloning either longer sequences or the full length sequences; see Ausubel, et al., *supra*. Much can be done by informatics and many sequences can be clustered to include multiple sequences corresponding to a single gene, e.g., systems such as UniGene (see, http://www.ncbi.nlm.nih.gov/UniGene/).

Once a lung cancer nucleic acid is identified, it can be cloned and, if necessary, its constituent parts recombined to form the entire lung cancer nucleic acid coding regions or the entire mRNA sequence. Once isolated from its natural source, e.g., contained within a plasmid or other vector or excised therefrom as a linear nucleic acid segment, the recombinant lung cancer nucleic acid can be further-used as a probe to identify and isolate

other lung cancer nucleic acids, e.g., extended coding regions. It can also be used as a "precursor" nucleic acid to make modified or variant lung cancer nucleic acids and proteins.

The lung cancer nucleic acids of the present invention are used in several ways. In a first embodiment, nucleic acid probes to the lung cancer nucleic acids are made and attached to biochips to be used in screening and diagnostic methods, as outlined below, or for administration, e.g., for gene therapy, RNAi, vaccine, and/or antisense applications. Alternatively, the lung cancer nucleic acids that include coding regions of lung cancer proteins can be put into expression vectors for the expression of lung cancer proteins, again for screening purposes or for administration to a patient.

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In a preferred embodiment, nucleic acid probes to lung cancer nucleic acids (both the nucleic acid sequences outlined in the figures and/or the complements thereof) are made. The nucleic acid probes attached to the biochip are designed to be substantially complementary to the lung cancer nucleic acids, i.e., the target sequence (either the target sequence of the sample or to other probe sequences, e.g., in sandwich assays), such that hybridization of the target sequence and the probes of the present invention occurs. As outlined below, this complementarity need not be perfect; there may be any number of base pair mismatches which will interfere with hybridization between the target sequence and the single stranded nucleic acids of the present invention. However, if the number of mutations is so great that no hybridization can occur under even the least stringent of hybridization conditions, the sequence is not a complementary target sequence. Thus, by "substantially complementary" herein is meant that the probes are sufficiently complementary to the target sequences to hybridize under appropriate reaction conditions, particularly high stringency conditions, as outlined herein.

A nucleic acid probe is generally single stranded but can be partially single and partially double stranded. The strandedness of the probe is dictated by the structure, composition, and properties of the target sequence. In general, the nucleic acid probes range from about 8 to about 100 bases long, with from about 10 to about 80 bases being preferred, and from about 30 to about 50 bases being particularly preferred. That is, generally complements of ORFs or whole genes are not used. In some embodiments, nucleic acids of lengths up to hundreds of bases can be used.

In a preferred embodiment, more than one probe per sequence is used, with either overlapping probes or probes to different sections of the target being used. That is, two, three, four or more probes, with three being preferred, are used to build in a redundancy for a

WO 02/086443 PCT/US02/12476 particular target. The probes can be overlapping (i.e., have some sequence in common), or

separate. In some cases, PCR primers may be used to amplify signal for higher sensitivity.

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As will be appreciated by those in the art, nucleic acids can be attached or immobilized to a solid support in a wide variety of ways. By "immobilized" and grammatical equivalents herein is meant the association or binding between the nucleic acid probe and the solid support is sufficient to be stable under the conditions of binding, washing, analysis, and removal as outlined below. The binding can typically be covalent or non-covalent. By "non-covalent binding" and grammatical equivalents herein is typically meant one or more of electrostatic, hydrophilic, and hydrophobic interactions. Included in non-covalent binding is the covalent attachment of a molecule, such as, streptavidin to the support and the non-covalent binding of the biotinylated probe to the streptavidin. By "covalent binding" and grammatical equivalents herein is meant that the two moieties, the solid support and the probe, are attached by at least one bond, including sigma bonds, pi bonds and coordination bonds. Covalent bonds can be formed directly between the probe and the solid support or can be formed by a cross linker or by inclusion of a specific reactive group on either the solid support or the probe or both molecules. Immobilization may also involve a combination of covalent and non-covalent interactions.

In general, the probes are attached to a biochip in a wide variety of ways, as will be appreciated by those in the art. As described herein, the nucleic acids can either be synthesized first, with subsequent attachment to the biochip, or can be directly synthesized on the biochip.

The biochip comprises a suitable solid substrate. By "substrate" or "solid support" or other grammatical equivalents herein is meant a material that can be modified for the attachment or association of the nucleic acid probes and is amenable to at least one detection method. Often the substrate may contain discrete individual sites appropriate for ndivitual partitioning and identification. As will be appreciated by those in the art, the number of possible substrates are very large, and include, but are not limited to, glass and modified or functionalized glass, plastics (including acrylics, polystyrene and copolymers of styrene and other materials, polypropylene, polyethylene, polybutylene, polyurethanes, Teflon, etc.), polysaccharides, nylon or nitrocellulose, resins, silica or silica-based materials including silicon and modified silicon, carbon, metals, inorganic glasses, plastics, etc. In general, the substrates allow optical detection and do not appreciably fluoresce. A preferred substrate is described in US application entitled Reusable Low Fluorescent Plastic Biochip, U.S.

Application Serial No. 09/270,214, filed March 15, 1999, herein incorporated by reference in its entirety.

Generally the substrate is planar, although as will be appreciated by those in the art, other configurations of substrates may be used as well. For example, the probes may be placed on the inside surface of a tube, for flow-through sample analysis to minimize sample volume. Similarly, the substrate may be flexible, such as a flexible foam, including closed cell foams made of particular plastics.

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In a preferred embodiment, the surface of the biochip and the probe may be derivatized with chemical functional groups for subsequent attachment of the two. Thus, e.g., the biochip is derivatized with a chemical functional group including, but not limited to, amino groups, carboxy groups, oxo groups and thiol groups, with amino groups being particularly preferred. Using these functional groups, the probes can be attached using functional groups on the probes. For example, nucleic acids containing amino groups can be attached to surfaces comprising amino groups, e.g., using linkers as are known in the art; e.g., homo-or hetero-bifunctional linkers as are well known (see 1994 Pierce Chemical Company catalog, technical section on cross-linkers, pages 155-200). In addition, in some cases, additional linkers, such as alkyl groups (including substituted and heteroalkyl groups) may be used.

In this embodiment, oligonucleotides are synthesized, and then attached to the surface of the solid support. Either the 5' or 3' terminus may be attached to the solid support, or attachment may be via linkage to an internal nucleoside.

In another embodiment, the immobilization to the solid support may be very strong, yet non-covalent. For example, biotinylated oligonucleotides can be made, which bind to surfaces covalently coated with streptavidin, resulting in attachment.

Alternatively, the oligonucleotides may be synthesized on the surface, as is known in the art. For example, photoactivation techniques utilizing photopolymerization compounds and techniques are used. In a preferred embodiment, the nucleic acids can be synthesized in situ, using known photolithographic techniques, such as those described in WO 95/25116; WO 95/35505; U.S. Patent Nos. 5,700,637 and 5,445,934; and references cited within, all of which are expressly incorporated by reference; these methods of attachment form the basis of the Affymetrix GeneChipTM technology.

Often, amplification-based assays are performed to measure the expression level of lung cancer-associated sequences. These assays are typically performed in conjunction with

reverse transcription. In such assays, a lung cancer-associated nucleic acid sequence acts as a template in an amplification reaction (e.g., Polymerase Chain Reaction, or PCR). In a quantitative amplification, the amount of amplification product will be proportional to the amount of template in the original sample. Comparison to appropriate controls provides a measure of the amount of lung cancer-associated RNA. Methods of quantitative amplification are well known to those of skill in the art. Detailed protocols for quantitative PCR are provided, e.g., in Innis, et al. (1990) PCR Protocols, A Guide to Methods and Applications.

In some embodiments, a TaqMan based assay is used to measure expression. TaqMan based assays use a fluorogenic oligonucleotide probe that contains a 5' fluorescent dye and a 3' quenching agent. The probe hybridizes to a PCR product, but cannot itself be extended due to a blocking agent at the 3' end. When the PCR product is amplified in subsequent cycles, the 5' nuclease activity of the polymerase, e.g., AmpliTaq, results in the cleavage of the TaqMan probe. This cleavage separates the 5' fluorescent dye and the 3' quenching agent, thereby resulting in an increase in fluorescence as a function of amplification (see, e.g., literature provided by Perkin-Elmer, e.g., www2.perkin-elmer.com).

Other suitable amplification methods include, but are not limited to, ligase chain reaction (LCR) (see Wu and Wallace (1989) Genomics 4:560, Landegren, et al. (1988) Science 241:1077, and Barringer, et al. (1990) Gene 89:117), transcription amplification (Kwoh, et al. (1989) Proc. Natl. Acad. Sci. USA 86:1173), self-sustained sequence replication (Guatelli, et al. (1990) Proc. Nat. Acad. Sci. USA 87:1874), dot PCR, and linker adapter PCR, etc.

Expression of lung cancer proteins from nucleic acids

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In a preferred embodiment, lung cancer nucleic acids, e.g., encoding lung cancer proteins, are used to make a variety of expression vectors to express lung cancer proteins which can then be used in screening assays, as described below. Expression vectors and recombinant DNA technology are well known to those of skill in the art (see, e.g., Ausubel, supra, and Fernandez and Hoeffler (eds 1999) Gene Expression Systems) and are used to express proteins. The expression vectors may be either self-replicating extrachromosomal vectors or vectors which integrate into a host genome. Generally, these expression vectors include transcriptional and translational regulatory nucleic acid operably linked to the nucleic acid encoding the lung cancer protein. The term "control sequences" refers to DNA

sequences used for the expression of an operably linked coding sequence in a particular host organism. Control sequences that are suitable for prokaryotes, e.g., include a promoter, optionally an operator sequence, and a ribosome binding site. Eukaryotic cells are known to utilize promoters, polyadenylation signals, and enhancers.

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Nucleic acid is "operably linked" when it is placed into a functional relationship with another nucleic acid sequence. For example, DNA for a presequence or secretory leader is operably linked to DNA for a polypeptide if it is expressed as a preprotein that participates in the secretion of the polypeptide; a promoter or enhancer is operably linked to a coding sequence if it affects the transcription of the sequence; or a ribosome binding site is operably linked to a coding sequence if it is positioned so as to facilitate translation. Generally, "operably linked" means that the DNA sequences being linked are contiguous, and, in the case of a secretory leader, contiguous and in reading phase. However, enhancers do not have to be contiguous. Linking is typically accomplished by ligation at convenient restriction sites. If such sites do not exist, synthetic oligonucleotide adaptors or linkers are used in accordance with conventional practice. Transcriptional and translational regulatory nucleic acid will generally be appropriate to the host cell used to express the lung cancer protein. Numerous types of appropriate expression vectors, and suitable regulatory sequences are known in the art for a variety of host cells.

In general, transcriptional and translational regulatory sequences may include, but are not limited to, promoter sequences, ribosomal binding sites, transcriptional start and stop sequences, translational start and stop sequences, and enhancer or activator sequences. In a preferred embodiment, the regulatory sequences include a promoter and transcriptional start and stop sequences.

Promoter sequences may be either constitutive or inducible promoters. The promoters may be either naturally occurring promoters or hybrid promoters. Hybrid promoters, which combine elements of more than one promoter, are also known in the art, and are useful in the present invention.

In addition, an expression vector may comprise additional elements. For example, the expression vector may have two replication systems, thus allowing it to be maintained in two organisms, e.g., in mammalian or insect cells for expression and in a prokaryotic host for cloning and amplification. Furthermore, for integrating expression vectors, the expression vector often contains at least one sequence homologous to the host cell genome, and preferably two homologous sequences which flank the expression construct. The integrating

vector may be directed to a specific locus in the host cell by selecting the appropriate homologous sequence for inclusion in the vector. Constructs for integrating vectors are well known in the art (e.g., Fernandez and Hoeffler, *supra*).

In addition, in a preferred embodiment, the expression vector contains a selectable marker gene to allow the selection of transformed host cells. Selection genes are well known in the art and will vary with the host cell used.

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The lung cancer proteins of the present invention are usually produced by culturing a host cell transformed with an expression vector containing nucleic acid encoding a lung cancer protein, under the appropriate conditions to induce or cause expression of the lung cancer protein. Conditions appropriate for lung cancer protein expression will vary with the choice of the expression vector and the host cell, and will be easily ascertained by one skilled in the art through routine experimentation or optimization. For example, the use of constitutive promoters in the expression vector will require optimizing the growth and proliferation of the host cell, while the use of an inducible promoter requires the appropriate growth conditions for induction. In addition, in some embodiments, the timing of the harvest is important. For example, the baculoviral systems used in insect cell expression are lytic viruses, and thus harvest time selection can be crucial for product yield.

Appropriate host cells include yeast, bacteria, archaebacteria, fungi, and insect and animal cells, including mammalian cells. Of particular interest are *Saccharomyces cerevisiae* and other yeasts, *E. coli*, *Bacillus subtilis*, Sf9 cells, C129 cells, 293 cells, *Neurospora*, BHK, CHO, COS, HeLa cells, HUVEC (human umbilical vein endothelial cells), THP1 cells (a macrophage cell line) and various other human cells and cell lines.

In a preferred embodiment, the lung cancer proteins are expressed in mammalian cells. Mammalian expression systems are also known in the art, and include retroviral and adenoviral systems. Of particular use as mammalian promoters are the promoters from mammalian viral genes, since the viral genes are often highly expressed and have a broad host range. Examples include the SV40 early promoter, mouse mammary tumor virus LTR promoter, adenovirus major late promoter, herpes simplex virus promoter, and the CMV promoter (see, e.g., Fernandez and Hoeffler, *supra*). Typically, transcription termination and polyadenylation sequences recognized by mammalian cells are regulatory regions located 3' to the translation stop codon and thus, together with the promoter elements, flank the coding sequence. Examples of transcription terminator and polyadenylation signals include those derived form SV40.

The methods of introducing exogenous nucleic acid into mammalian hosts, as well as other hosts, is well known in the art, and will vary with the host cell used. Techniques include dextran-mediated transfection, calcium phosphate precipitation, polybrene mediated transfection, protoplast fusion, electroporation, viral infection, encapsulation of the polynucleotide(s) in liposomes, and direct microinjection of the DNA into nuclei.

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In a preferred embodiment, lung cancer proteins are expressed in bacterial systems. Promoters from bacteriophage may also be used and are known in the art. In addition, synthetic promoters and hybrid promoters are also useful; e.g., the tac promoter is a hybrid of the trp and lac promoter sequences. Furthermore, a bacterial promoter can include naturally occurring promoters of non-bacterial origin that have the ability to bind bacterial RNA polymerase and initiate transcription. In addition to a functioning promoter sequence, an efficient ribosome binding site is desirable. The expression vector may also include a signal peptide sequence that provides for secretion of the lung cancer protein in bacteria. The protein is either secreted into the growth media (gram-positive bacteria) or into the periplasmic space, located between the inner and outer membrane of the cell (gram-negative bacteria). The bacterial expression vector may also include a selectable marker gene to allow for the selection of bacterial strains that have been transformed. Suitable selection genes include genes which render the bacteria resistant to drugs such as ampicillin, chloramphenicol, erythromycin, kanamycin, neomycin and tetracycline. Selectable markers also include biosynthetic genes, such as those in the histidine, tryptophan and leucine biosynthetic pathways. These components are assembled into expression vectors. Expression vectors for bacteria are well known in the art, and include vectors for Bacillus subtilis, E. coli, Streptococcus cremoris, and Streptococcus lividans, among others (e.g., Fernandez and Hoeffler, supra). The bacterial expression vectors are transformed into bacterial host cells using techniques well known in the art, such as calcium chloride treatment, electroporation, and others.

In one embodiment, lung cancer proteins are produced in insect cells. Expression vectors for the transformation of insect cells, and in particular, baculovirus-based expression vectors, are well known in the art.

In a preferred embodiment, lung cancer protein is produced in yeast cells. Yeast expression systems are well known in the art, and include expression vectors for Saccharomyces cerevisiae, Candida albicans and C. maltosa, Hansenula polymorpha,

Kluyveromyces fragilis and K. lactis, Pichia guillerimondii, and P. pastoris, Schizosaccharomyces pombe, and Yarrowia lipolytica.

The lung cancer protein may also be made as a fusion protein, using techniques well known in the art. Thus, e.g., for the creation of monoclonal antibodies, if the desired epitope is small, the lung cancer protein may be fused to a carrier protein to form an immunogen. Alternatively, the lung cancer protein may be made as a fusion protein to increase expression for affinity purification purposes, or for other reasons. For example, when the lung cancer protein is a lung cancer peptide, the nucleic acid encoding the peptide may be linked to other nucleic acid for expression purposes.

In a preferred embodiment, the lung cancer protein is purified or isolated after expression. Lung cancer proteins may be isolated or purified in a variety of appropriate ways. Standard purification methods include electrophoretic, molecular, immunological and chromatographic techniques, including ion exchange, hydrophobic, affinity, and reverse-phase HPLC chromatography, and chromatofocusing. For example, the lung cancer protein may be purified using a standard anti-lung cancer protein antibody column. Ultrafiltration and diafiltration techniques, in conjunction with protein concentration, are also useful. For general guidance in suitable purification techniques, see Scopes (1982) Protein Purification. The degree of purification necessary will vary depending on the use of the lung cancer protein. In some instances no purification will be necessary.

Once expressed and purified if necessary, the lung cancer proteins and nucleic acids are useful in a number of applications. They may be used as immunoselection reagents, as vaccine reagents, as screening agents, therapeutic entities, for production of antibodies, as transcription or translation inhibitors, etc.

25 Variants of lung cancer proteins

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In one embodiment, the lung cancer proteins are derivative or variant lung cancer proteins as compared to the wild-type sequence. That is, as outlined more fully below, the derivative lung cancer peptide will often contain at least one amino acid substitution, deletion or insertion, with amino acid substitutions being particularly preferred. The amino acid substitution, insertion or deletion may occur at a particular residue within the lung cancer peptide.

Also included within one embodiment of lung cancer proteins of the present invention are amino acid sequence variants. These variants typically fall into one or more of three

classes: substitutional, insertional or deletional variants. These variants ordinarily are prepared by site specific mutagenesis of nucleotides in the DNA encoding the lung cancer protein, using cassette or PCR mutagenesis or other techniques, to produce DNA encoding the variant, and thereafter expressing the DNA in recombinant cell culture as outlined above. However, variant lung cancer protein fragments having up to about 100-150 residues may be prepared by *in vitro* synthesis. Amino acid sequence variants are characterized by the predetermined nature of the variation, a feature that sets them apart from naturally occurring allelic or interspecies variation of the lung cancer protein amino acid sequence. The variants typically exhibit a similar qualitative biological activity as the naturally occurring analogue, although variants can also be selected which have modified characteristics as will be more fully outlined below.

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While the site or region for introducing an amino acid sequence variation is often predetermined, the mutation per se need not be predetermined. For example, in order to optimize the performance of a mutation at a given site, random mutagenesis may be conducted at the target codon or region and the expressed lung cancer variants screened for the optimal combination of desired activity. Techniques exist for making substitution mutations at predetermined sites in DNA having a known sequence, e.g., M13 primer mutagenesis and PCR mutagenesis. Screening of mutants is often done using assays of lung cancer protein activities.

Amino acid substitutions are typically of single residues; insertions usually will be on the order of from about 1 to 20 amino acids, although considerably larger insertions may be occasionally tolerated. Deletions generally range from about 1 to about 20 residues, although in some cases deletions may be much larger.

Substitutions, deletions, insertions or any combination thereof may be used to arrive at a final derivative. Generally these changes are done on a few amino acids to minimize the alteration of the molecule. Larger changes may be tolerated in certain circumstances. When small alterations in the characteristics of a lung cancer protein are desired, substitutions are generally made in accordance with the amino acid substitution chart provided in the definition section.

Variants typically exhibit essentially the same qualitative biological activity and will elicit the same immune response as a naturally-occurring analog, although variants also are selected to modify the characteristics of lung cancer proteins as needed. Alternatively, the

variant may be designed or reorganized such that a biological activity of the lung cancer protein is altered. For example, glycosylation sites may be added, altered, or removed.

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Covalent modifications of lung cancer polypeptides are included within the scope of this invention. One type of covalent modification includes reacting targeted amino acid residues of a lung cancer polypeptide with an organic derivatizing agent that is capable of reacting with selected side chains or the N-or C-terminal residues of a lung cancer polypeptide. Derivatization with bifunctional agents is useful, for instance, for crosslinking lung cancer polypeptides to a water-insoluble support matrix or surface for use in a method for purifying anti-lung cancer polypeptide antibodies or screening assays, as is more fully described below. Commonly used crosslinking agents include, e.g., 1,1-bis(diazoacetyl)-2-phenylethane, glutaraldehyde, N-hydroxysuccinimide esters, e.g., esters with 4-azidosalicylic acid, homobifunctional imidoesters, including disuccinimidyl esters such as 3,3'-dithiobis(succinimidylpropionate), bifunctional maleimides such as bis-N-maleimido-1,8-octane and agents such as methyl-3-((p-azidophenyl)dithio)propioimidate.

Other modifications include deamidation of glutaminyl and asparaginyl residues to the corresponding glutamyl and aspartyl residues, respectively, hydroxylation of proline and lysine, phosphorylation of hydroxyl groups of serinyl, threonyl or tyrosyl residues, methylation of the γ-amino groups of lysine, arginine, and histidine side chains (Creighton (1983) Proteins: Structure and Molecular Properties, pp. 79-86), acetylation of the N-terminal amine, and amidation of any C-terminal carboxyl group.

Another type of covalent modification of the lung cancer polypeptide encompassed by this invention is an altered native glycosylation pattern of the polypeptide. "Altering the native glycosylation pattern" is intended herein to mean adding to or deleting one or more carbohydrate moieties of a native sequence lung cancer polypeptide. Glycosylation patterns can be altered in many ways. For example the use of different cell types to express lung cancer-associated sequences can result in different glycosylation patterns.

Addition of glycosylation sites to lung cancer polypeptides may also be accomplished by altering the amino acid sequence thereof. The alteration may be made, e.g., by the addition of, or substitution by, one or more serine or threonine residues to the native sequence lung cancer polypeptide (for O-linked glycosylation sites). The lung cancer amino acid sequence may optionally be altered through changes at the DNA level, particularly by mutating the DNA encoding the lung cancer polypeptide at preselected bases such that codons are generated that will translate into the desired amino acids.

Another means of increasing the number of carbohydrate moieties on the lung cancer polypeptide is by chemical or enzymatic coupling of glycosides to the polypeptide. Such methods are described in the art, e.g., in WO 87/05330, and in Aplin and Wriston (1981) CRC Crit. Rev. Biochem., pp. 259-306.

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Removal of carbohydrate moieties present on the lung cancer polypeptide may be accomplished chemically or enzymatically or by mutational substitution of codons encoding for amino acid residues that serve as targets for glycosylation. Chemical deglycosylation techniques are known in the art and described, for instance, by Hakimuddin, et al. (1987)

Arch. Biochem. Biophys., 259:52 and by Edge, et al. (1981) Anal. Biochem., 118:131.

Enzymatic cleavage of carbohydrate moieties on polypeptides can be achieved by the use of a variety of endo-and exo-glycosidases as described by Thotakura, et al. (1987) Meth.

Enzymol., 138:350.

Another type of covalent modification of lung cancer comprises linking the lung cancer polypeptide to one of a variety of nonproteinaceous polymers, e.g., polyethylene glycol, polypropylene glycol, or polyoxyalkylenes, in the manner set forth in U.S. Patent Nos. 4,640,835; 4,496,689; 4,301,144; 4,670,417; 4,791,192, or 4,179,337.

Lung cancer polypeptides of the present invention may also be modified in a way to form chimeric molecules comprising a lung cancer polypeptide fused to another, heterologous polypeptide or amino acid sequence. In one embodiment, such a chimeric molecule comprises a fusion of a lung cancer polypeptide with a tag polypeptide which provides an epitope to which an anti-tag antibody can selectively bind. The epitope tag is generally placed at the amino-or carboxyl-terminus of the lung cancer polypeptide. The presence of such epitope-tagged forms of a lung cancer polypeptide can be detected using an antibody against the tag polypeptide. Also, provision of the epitope tag enables the lung cancer polypeptide to be readily purified by affinity purification using an anti-tag antibody or another type of affinity matrix that binds to the epitope tag. In an alternative embodiment, the chimeric molecule may comprise a fusion of a lung cancer polypeptide with an immunoglobulin or a particular region of an immunoglobulin. For a bivalent form of the chimeric molecule, such a fusion could be to the Fc region of an IgG molecule.

Various tag polypeptides and their respective antibodies are well known and examples include poly-histidine (poly-his) or poly-histidine-glycine (poly-his-gly) tags; HIS6 and metal chelation tags, the flu HA tag polypeptide and its antibody 12CA5 (Field, et al. (1988) Mol. Cell. Biol. 8:2159-2165); the c-myc tag and the 8F9, 3C7, 6E10, G4, B7 and 9E10 antibodies

thereto (Evan, et al. (1985) Molecular and Cellular Biology 5:3610-3616); and the Herpes Simplex virus glycoprotein D (gD) tag and its antibody (Paborsky, et al. (1990) Protein Engineering 3(6):547-553). Other tag polypeptides include the Flag-peptide (Hopp, et al. (1988) BioTechnology 6:1204-1210); the KT3 epitope peptide (Martin, et al. (1992) Science 255:192-194); tubulin epitope peptide (Skinner, et al. (1991) J. Biol. Chem. 266:15163-15166); and the T7 gene 10 protein peptide tag (Lutz-Freyermuth, et al. (1990) Proc. Nat'l Acad. Sci. USA 87:6393-6397).

Also included are other lung cancer proteins of the lung cancer family, and lung cancer proteins from other organisms, which are cloned and expressed as outlined below. Thus, probe or degenerate polymerase chain reaction (PCR) primer sequences may be used to find other related lung cancer proteins from primates or other organisms. As will be appreciated by those in the art, particularly useful probe and/or PCR primer sequences include unique areas of the lung cancer nucleic acid sequence. As is generally known in the art, preferred PCR primers are from about 15 to about 35 nucleotides in length, with from about 20 to about 30 being preferred, and may contain inosine as needed. PCR reaction conditions are well known in the art (e.g., Innis, PCR Protocols, *supra*).

Antibodies to lung cancer proteins

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In a preferred embodiment, when a lung cancer protein is to be used to generate antibodies, e.g., for immunotherapy or immunodiagnosis, the lung cancer protein should share at least one epitope or determinant with the full length protein. By "epitope" or "determinant" herein is typically meant a portion of a protein which will generate and/or bind an antibody or T-cell receptor in the context of MHC. Thus, in most instances, antibodies made to a smaller lung cancer protein will be able to bind to the full-length protein, particularly linear epitopes. In a preferred embodiment, the epitope is unique; that is, antibodies generated to a unique epitope show little or no cross-reactivity.

Methods of preparing polyclonal antibodies are well known (e.g., Coligan, supra; and Harlow and Lane, supra). Polyclonal antibodies can be raised in a mammal, e.g., by one or more injections of an immunizing agent and, if desired, an adjuvant. Typically, the immunizing agent and/or adjuvant will be injected in the mammal by multiple subcutaneous or intraperitoneal injections. The immunizing agent may include a protein encoded by a nucleic acid of Tables 1A-16 or fragment thereof or a fusion protein thereof. It may be useful to conjugate the immunizing agent to a protein known to be immunogenic in the mammal

being immunized. Immunogenic proteins include, e.g., keyhole limpet hemocyanin, serum albumin, bovine thyroglobulin, and soybean trypsin inhibitor. Adjuvants include, e.g., Freund's complete adjuvant and MPL-TDM adjuvant (monophosphoryl Lipid A, synthetic trehalose dicorynomycolate). The immunization protocol may be selected by one skilled in the art.

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The antibodies may, alternatively, be monoclonal antibodies. Monoclonal antibodies may be prepared using hybridoma methods, such as those described by Kohler and Milstein (1975) Nature 256:495. In a hybridoma method, a mouse, hamster, or other appropriate host animal, is typically immunized with an immunizing agent to elicit lymphocytes that produce or are capable of producing antibodies that will specifically bind to the immunizing agent. Alternatively, the lymphocytes may be immunized in vitro. The immunizing agent will typically include a polypeptide encoded by a nucleic acid of the tables, or fragment thereof, or a fusion protein thereof. Generally, either peripheral blood lymphocytes ("PBLs") are used if cells of human origin are desired, or spleen cells or lymph node cells are used if nonhuman mammalian sources are desired. The lymphocytes are then fused with an immortalized cell line using a suitable fusing agent, such as polyethylene glycol, to form a hybridoma cell (Goding (1986) Monoclonal Antibodies: Principles and Practice, pp. 59-103). Immortalized cell lines are usually transformed mammalian cells, particularly myeloma cells of rodent, bovin, or primate origin. Usually, rat or mouse myeloma cell lines are employed. The hybridoma cells may be cultured in a suitable culture medium that preferably contains one or more substances that inhibit the growth or survival of the unfused, immortalized cells. For example, if the parental cells lack the enzyme hypoxanthine guanine phosphoribosyl transferase (HGPRT or HPRT), the culture medium for the hybridomas typically will include hypoxanthine, aminopterin, and thymidine ("HAT medium"), which substances prevent the growth of HGPRT-deficient cells.

In one embodiment, the antibodies are bispecific antibodies. Bispecific antibodies are typically monoclonal, preferably human or humanized, antibodies that have binding specificities for at least two different antigens or that have binding specificities for two epitopes on the same antigen. In one embodiment, one of the binding specificities is for a protein encoded by a nucleic acid of the tables or a fragment thereof, the other one is for any other antigen, and preferably for a cell-surface protein or receptor or receptor subunit, preferably one that is tumor specific. Alternatively, tetramer-type technology may create multivalent reagents.

In a preferred embodiment, the antibodies to lung cancer protein are capable of reducing or eliminating a biological function of a lung cancer protein, in a naked form or conjugated to an effector moiety. That is, the addition of anti-lung cancer protein antibodies (either polyclonal or preferably monoclonal) to lung cancer tissue (or cells containing lung cancer) may reduce or eliminate the lung cancer. Generally, at least a 25% decrease in activity, growth, size or the like is preferred, with at least about 50% being particularly preferred and about a 95-100% decrease being especially preferred.

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In a preferred embodiment the antibodies to the lung cancer proteins are humanized antibodies (e.g., Xenerex Biosciences, Medarex, Inc., Abgenix, Inc., Protein Design Labs, Inc.) Humanized forms of non-human (e.g., murine) antibodies are chimeric molecules of immunoglobulins, immunoglobulin chains or fragments thereof (such as Fv, Fab, Fab', F(ab')2 or other antigen-binding subsequences of antibodies) which contain minimal sequence derived from non-human immunoglobulin. Humanized antibodies include human immunoglobulins (recipient antibody) in which residues from a complementary determining region (CDR) of the recipient are replaced by residues from a CDR of a non-human species (donor antibody) such as mouse, rat or rabbit having the desired specificity, affinity and capacity. In some instances, Fv framework residues of a human immunoglobulin are replaced by corresponding non-human residues. Humanized antibodies may also comprise residues which are found neither in the recipient antibody nor in the imported CDR or framework sequences. In general, a humanized antibody will comprise substantially all of at least one, and typically two, variable domains, in which all or substantially all of the CDR regions correspond to those of a non-human immunoglobulin and all or substantially all of the framework (FR) regions are those of a human immunoglobulin consensus sequence. A humanized antibody optimally also will typically comprise at least a portion of an immunoglobulin constant region (Fc), typically that of a human immunoglobulin (Jones, et al. (1986) Nature 321:522-525; Riechmann, et al. (1988) Nature 332:323-329; and Presta (1992) Curr. Op. Struct. Biol. 2:593-596). Humanization can be performed following the method of Winter and co-workers (Jones, et al. (1986) Nature 321:522-525; Riechmann, et al. (1988) Nature 332:323-327; Verhoeyen, et al. (1988) Science 239:1534-1536), by substituting rodent CDRs or CDR sequences for corresponding sequences of a human antibody. Accordingly, such humanized antibodies are chimeric antibodies (U.S. Patent No. 4,816,567), wherein substantially less than an intact human variable domain has been substituted by corresponding sequence from a non-human species.

Human-like antibodies can also be produced using various techniques known in the art, including phage display libraries (Hoogenboom and Winter (1991) J. Mol. Biol. 227:381; Marks, et al. (1991) J. Mol. Biol. 222:581). The techniques of Cole, et al. and Boerner, et al. are also available for the preparation of human monoclonal antibodies (Cole, et al. (1985) Monoclonal Antibodies and Cancer Therapy, p. 77 and Boerner, et al. (1991) J. Immunol. 147(1):86-95). Similarly, human antibodies can be made by introducing human immunoglobulin loci into transgenic animals, e.g., mice in which the endogenous immunoglobulin genes have been partially or completely inactivated. Upon challenge, human antibody production is observed, which closely resembles that seen in humans in nearly all respects, including gene rearrangement, assembly, and antibody repertoire. This approach is described, e.g., in U.S. Patent Nos. 5,545,807; 5,545,806; 5,569,825; 5,625,126; 5.633.425; 5.661.016, and in the following scientific publications: Marks, et al. (1992) Bio/Technology 10:779-783; Lonberg, et al. (1994) Nature 368:856-859; Morrison (1994) Nature 368:812-13; Fishwild, et al. (1996) Nature Biotechnology 14:845-51; Neuberger (1996) Nature Biotechnology 14:826; and Lonberg and Huszar (1995) Intern. Rev. Immunol. 13:65-93.

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By immunotherapy is meant treatment of lung cancer with an antibody raised against a lung cancer proteins. As used herein, immunotherapy can be passive or active. Passive immunotherapy as defined herein is the passive transfer of antibody to a recipient (patient). Active immunization is the induction of antibody and/or T-cell responses in a recipient (patient). Induction of an immune response is the result of providing the recipient with an antigen to which antibodies are raised. The antigen may be provided by injecting a polypeptide against which antibodies are desired to be raised into a recipient, or contacting the recipient with a nucleic acid capable of expressing the antigen and under conditions for expression of the antigen, leading to an immune response.

In a preferred embodiment the lung cancer proteins against which antibodies are raised are secreted proteins as described above. Without being bound by theory, antibodies used for treatment, may bind and prevent the secreted protein from binding to its receptor, thereby inactivating the secreted lung cancer protein.

In another preferred embodiment, the lung cancer protein to which antibodies are raised is a transmembrane protein. Without being bound by theory, antibodies used for treatment may bind the extracellular domain of the lung cancer protein and prevent it from binding to other proteins, such as circulating ligands or cell-associated molecules. The

antibody may cause down-regulation of the transmembrane lung cancer protein. The antibody may be a competitive, non-competitive or uncompetitive inhibitor of protein binding to the extracellular domain of the lung cancer protein. The antibody may be an antagonist of the lung cancer protein or may prevent activation of a transmembrane lung cancer protein, or may induce or suppress a particular cellular pathway. In some embodiments, when the antibody prevents the binding of other molecules to the lung cancer protein, the antibody prevents growth of the cell. The antibody may also be used to target or sensitize the cell to cytotoxic agents, including, but not limited to TNF-α, TNF-β, IL-1, INF-γ, and IL-2, or chemotherapeutic agents including 5FU, vinblastine, actinomycin D, cisplatin, methotrexate, and the like. In some instances the antibody may belong to a sub-type that activates serum complement when complexed with the transmembrane protein thereby mediating cytotoxicity or antigen-dependent cytotoxicity (ADCC). Thus, lung cancer may be treated by administering to a patient antibodies directed against the transmembrane lung cancer protein. Antibody-labeling may activate a co-toxin, localize a toxin payload, or otherwise provide means to locally ablate cells.

In another preferred embodiment, the antibody is conjugated to an effector moiety. The effector moiety can be various molecules, including labeling moieties such as radioactive labels or fluorescent labels, or can be a therapeutic moiety. In one aspect the therapeutic moiety is a small molecule that modulates the activity of a lung cancer protein. In another aspect the therapeutic moiety may modulate an activity of molecules associated with or in close proximity to a lung cancer protein. The therapeutic moiety may inhibit enzymatic or signaling activity such as protease or collagenase activity associated with lung cancer.

In a preferred embodiment, the therapeutic moiety can also be a cytotoxic agent. In this method, targeting the cytotoxic agent to lung cancer tissue or cells results in a reduction in the number of afflicted cells, thereby reducing symptoms associated with lung cancer. Cytotoxic agents are numerous and varied and include, but are not limited to, cytotoxic drugs or toxins or active fragments of such toxins. Suitable toxins and their corresponding fragments include diphtheria A chain, exotoxin A chain, ricin A chain, abrin A chain, curcin, crotin, phenomycin, enomycin, saporin, auristatin, and the like. Cytotoxic agents also include radiochemicals made by conjugating radioisotopes to antibodies raised against lung cancer proteins, or binding of a radionuclide to a chelating agent that has been covalently attached to the antibody. Targeting the therapeutic moiety to transmembrane lung cancer proteins not only serves to increase the local concentration of therapeutic moiety in the lung cancer

afflicted area, but also serves to reduce deleterious side effects that may be associated with the untargeted therapeutic moiety.

In another preferred embodiment, the lung cancer protein against which the antibodies are raised is an intracellular protein. In this case, the antibody may be conjugated to a protein or other entity which facilitates entry into the cell. In one case, the antibody enters the cell by endocytosis. In another embodiment, a nucleic acid encoding the antibody is administered to the individual or cell. Moreover, wherein the lung cancer protein can be targeted within a cell, i.e., the nucleus, an antibody theretomay contain a signal for that target localization, i.e., a nuclear localization signal.

The lung cancer antibodies of the invention specifically bind to lung cancer proteins. By "specifically bind" herein is meant that the antibodies bind to the protein with a K_d of at least about 0.1 mM, more usually at least about 1 μ M, preferably at least about 0.1 μ M or better, and most preferably, 0.01 μ M or better. Selectivity of binding to the specific target and not to related other sequences is also important.

Detection of lung cancer sequence for diagnostic and therapeutic applications

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In one aspect, the RNA expression levels of genes are determined for different cellular states in the lung cancer phenotype. Expression levels of genes in normal tissue (e.g., not undergoing lung cancer), in lung cancer tissue (and in some cases, for varying severities of lung cancer that relate to prognosis, as outlined below), or in non-malignant disease are evaluated to provide expression profiles. A gene expression profile of a particular cell state or point of development is essentially a "fingerprint" of the state of the cell. While two states may have a particular gene similarly expressed, the evaluation of a number of genes simultaneously allows the generation of a gene expression profile that is reflective of the state of the cell. By comparing expression profiles of cells in different states, information regarding which genes are important (including both up- and down-regulation of genes) in each of these states is obtained. Then, diagnosis may be performed or confirmed to determine whether a tissue sample has the gene expression profile of normal or cancerous tissue. This will provide for molecular diagnosis of related conditions.

"Differential expression," or grammatical equivalents as used herein, refers to qualitative or quantitative differences in the temporal and/or cellular gene expression patterns within and among cells and tissue. Thus, a differentially expressed gene can qualitatively have its expression altered, including an activation or inactivation, in, e.g.,

PCT/US02/12476 WO 02/086443 normal versus lung cancer tissue. Genes may be turned on or turned off in a particular state, relative to another state thus permitting comparison of two or more states. A qualitatively regulated gene will exhibit an expression pattern within a state or cell type which is detectable by standard techniques. Some genes will be expressed in one state or cell type, but not in both. Alternatively, the difference in expression may be quantitative, e.g., in that expression is increased or decreased; i.e., gene expression is either upregulated, resulting in an increased amount of transcript, or downregulated, resulting in a decreased amount of transcript. The degree to which expression differs need only be large enough to quantify via standard characterization techniques as outlined below, such as by use of Affymetrix GeneChip™ expression arrays, Lockhart (1996) Nature Biotechnology 14:1675-1680, hereby expressly incorporated by reference. Other techniques include, but are not limited to, quantitative reverse transcriptase PCR, northern analysis and RNase protection. As outlined above, preferably the change in expression (i.e., upregulation or downregulation) is typically at least about 50%, more preferably at least about 100%, more preferably at least about 150%, more preferably at least about 200%, with from 300 to at least 1000% being especially preferred.

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Evaluation may be at the gene transcript or the protein level. The amount of gene expression may be monitored using nucleic acid probes to the RNA or DNA equivalent of the gene transcript, and the quantification of gene expression levels, or, alternatively, the final gene product itself (protein) can be monitored, e.g., with antibodies to the lung cancer protein and standard immunoassays (ELISAs, etc.) or other techniques, including mass spectroscopy assays, 2D gel electrophoresis assays, etc. Proteins corresponding to lung cancer genes, e.g., those identified as being important in a lung cancer or disease phenotype, can be evaluated in a lung cancer diagnostic test. In a preferred embodiment, gene expression monitoring is performed simultaneously on a number of genes.

The lung cancer nucleic acid probes may be attached to biochips as outlined herein for the detection and quantification of lung cancer sequences in a particular cell. The assays are further described below in the example. PCR techniques can be used to provide greater sensitivity. Multiple protein expression monitoring can be performed as well. Similarly, these assays may be performed on an individual basis as well.

In a preferred embodiment nucleic acids encoding the lung cancer protein are detected. Although DNA or RNA encoding the lung cancer protein may be detected, of particular interest are methods wherein an mRNA encoding a lung cancer protein is detected.

Probes to detect mRNA can be a nucleotide/deoxynucleotide probe that is complementary to and hybridizes with the mRNA and includes, but is not limited to, oligonucleotides, cDNA or RNA. Probes also should contain a detectable label, as defined herein. In one method the mRNA is detected after immobilizing the nucleic acid to be examined on a solid support such as nylon membranes and hybridizing the probe with the sample. Following washing to remove the non-specifically bound probe, the label is detected. In another method detection of the mRNA is performed *in situ*. In this method permeabilized cells or tissue samples are contacted with a detectably labeled nucleic acid probe for sufficient time to allow the probe to hybridize with the target mRNA. Following washing to remove the non-specifically bound probe, the label is detected. For example a digoxygenin labeled riboprobe (RNA probe) that is complementary to the mRNA encoding a lung cancer protein is detected by binding the digoxygenin with an anti-digoxygenin secondary antibody and developed with nitro blue tetrazolium and 5-bromo-4-chloro-3-indoyl phosphate.

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In a preferred embodiment, various proteins from the three classes of proteins as described herein (secreted, transmembrane or intracellular proteins) are used in diagnostic assays. The lung cancer proteins, antibodies, nucleic acids, modified proteins and cells containing lung cancer sequences are used in diagnostic assays. This can be performed on an individual gene or corresponding polypeptide level. In a preferred embodiment, the expression profiles are used, preferably in conjunction with high throughput screening techniques to allow monitoring for expression profile genes and/or corresponding polypeptides.

As described and defined herein, lung cancer proteins, including intracellular, transmembrane, or secreted proteins, find use as markers of lung cancer, e.g., for prognostic or diagnostic purposes. Detection of these proteins in putative lung cancer tissue allows for detection, prognosis, or diagnosis of lung cancer or similar disease, and perhaps for selection of therapeutic strategy. In one embodiment, antibodies are used to detect lung cancer proteins. A preferred method separates proteins from a sample by electrophoresis on a gel (typically a denaturing and reducing protein gel, but may be another type of gel, including isoelectric focusing gels and the like). Following separation of proteins, the lung cancer protein is detected, e.g., by immunoblotting with antibodies raised against the lung cancer protein. Methods of immunoblotting are well known to those of ordinary skill in the art.

In another preferred method, antibodies to the lung cancer protein find use in *in situ* imaging techniques, e.g., in histology (e.g., Asai (ed. 1993) Methods in Cell Biology:

Antibodies in Cell Biology, volume 37. In this method cells are contacted with from one to many antibodies to the lung cancer protein(s). Following washing to remove non-specific antibody binding, the presence of the antibody or antibodies is detected. In one embodiment the antibody is detected by incubating with a secondary antibody that contains a detectable label, e.g., multicolor fluorescence or confocal imaging. In another method the primary antibody to the lung cancer protein(s) contains a detectable label, e.g., an enzyme marker that can act on a substrate. In another preferred embodiment each one of multiple primary antibodies contains a distinct and detectable label. This method finds particular use in simultaneous screening for a plurality of lung cancer proteins. Many other histological imaging techniques are also provided by the invention.

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In a preferred embodiment the label is detected in a fluorometer which has the ability to detect and distinguish emissions of different wavelengths. In addition, a fluorescence activated cell sorter (FACS) can be used in the method.

In another preferred embodiment, antibodies find use in diagnosing lung cancer from blood, serum, plasma, stool, and other samples. Such samples, therefore, are useful as samples to be probed or tested for the presence of lung cancer proteins. Antibodies can be used to detect a lung cancer protein by previously described immunoassay techniques including ELISA, immunoblotting (western blotting), immunoprecipitation, BIACORE technology and the like. Conversely, the presence of antibodies may indicate an immune response against an endogenous lung cancer protein or vaccine.

In a preferred embodiment, in situ hybridization of labeled lung cancer nucleic acid probes to tissue arrays is done. For example, arrays of tissue samples, including lung cancer tissue and/or normal tissue, are made. In situ hybridization (see, e.g., Ausubel, supra) is then performed. When comparing the fingerprints between an individual and a standard, the skilled artisan can make a diagnosis, a prognosis, or a prediction based on the findings. It is further understood that the genes which indicate the diagnosis may differ from those which indicate the prognosis and molecular profiling of the condition of the cells may lead to distinctions between responsive or refractory conditions or may be predictive of outcomes.

In a preferred embodiment, the lung cancer proteins, antibodies, nucleic acids, modified proteins and cells containing lung cancer sequences are used in prognosis assays. As above, gene expression profiles can be generated that correlate to lung cancer, clinical, pathological, or other information, in terms of long term prognosis. Again, this may be done on either a protein or gene level, with the use of genes being preferred. Single or multiple

genes may be useful in various combinations. As above, lung cancer probes may be attached to biochips for the detection and quantification of lung cancer sequences in a tissue or patient. The assays proceed as outlined above for diagnosis. PCR method may provide more sensitive and accurate quantification.

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Assays for therapeutic compounds

In a preferred embodiment, the proteins, nucleic acids, and antibodies as described herein are used in drug screening assays. The lung cancer proteins, antibodies, nucleic acids, modified proteins and cells containing lung cancer sequences are used in drug screening assays or by evaluating the effect of drug candidates on a "gene expression profile" or expression profile of polypeptides. In a preferred embodiment, the expression profiles are used, preferably in conjunction with high throughput screening techniques to allow monitoring for expression profile genes after treatment with a candidate agent (e.g., Zlokarnik, et al. (1998) Science 279:84-8; Heid (1996) Genome Res. 6:986-94.

In a preferred embodiment, the lung cancer proteins, antibodies, nucleic acids, modified proteins and cells containing the native or modified lung cancer proteins are used in screening assays. That is, the present invention provides novel methods for screening for compositions which modulate the lung cancer phenotype or an identified physiological function of a lung cancer protein. As above, this can be done on an individual gene level or by evaluating the effect of drug candidates on a "gene expression profile". In a preferred embodiment, the expression profiles are used, preferably in conjunction with high throughput screening techniques to allow monitoring for expression profile genes after treatment with a candidate agent, see Zlokarnik, *supra*.

Having identified differentially expressed genes herein, a variety of assays may be performed. In a preferred embodiment, assays may be run on an individual gene or protein level. That is, having identified a particular gene with altered regulation in lung cancer, test compounds can be screened for the ability to modulate gene expression or for binding to the lung cancer protein. "Modulation" thus includes an increase or a decrease in gene expression. The preferred amount of modulation will depend on the original change of the gene expression in normal versus tissue undergoing lung cancer, with changes of at least 10%, preferably 50%, more preferably 100-300%, and in some embodiments 300-1000% or greater. Thus, if a gene exhibits a 4-fold increase in lung cancer tissue compared to normal tissue, a decrease of about four-fold is often desired; similarly, a 10-fold decrease in lung

WO 02/086443 PCT/US02/12476 cancer tissue compared to normal tissue often provides a target value of a 10-fold increase in

expression to be induced by the test compound.

The amount of gene expression may be monitored using nucleic acid probes and the quantification of gene expression levels, or, alternatively, the gene product itself can be monitored, e.g., through the use of antibodies to the lung cancer protein and standard immunoassays. Proteomics and separation techniques may also allow quantification of expression.

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In a preferred embodiment, gene or protein expression monitoring of a number of entities, i.e., an expression profile, is monitored simultaneously. Such profiles will typically involve a plurality of those entities described herein.

In this embodiment, the lung cancer nucleic acid probes are attached to biochips as outlined herein for the detection and quantification of lung cancer sequences in a particular cell. Alternatively, PCR may be used. Thus, a series, e.g., of microtiter plate, may be used with dispensed primers in desired wells. A PCR reaction can then be performed and analyzed for each well.

Expression monitoring can be performed to identify compounds that modify the expression of one or more lung cancer-associated sequences, e.g., a polynucleotide sequence set out in the tables. Generally, in a preferred embodiment, a test compound is added to the cells prior to analysis. Moreover, screens are also provided to identify agents that modulate lung cancer, modulate lung cancer proteins, bind to a lung cancer protein, or interfere with the binding of a lung cancer protein and an antibody, substrate, or other binding partner.

The term "test compound" or "drug candidate" or "modulator" or grammatical equivalents as used herein describes a molecule, e.g., protein, oligopeptide, small organic molecule, polysaccharide, polynucleotide, etc., to be tested for the capacity to directly or indirectly alter the lung cancer phenotype or the expression of a lung cancer sequence, e.g., a nucleic acid or protein sequence. In preferred embodiments, modulators alter expression profiles of nucleic acids or proteins provided herein. In one embodiment, the modulator suppresses a lung cancer phenotype, e.g., to a normal or non-malignant tissue fingerprint. In another embodiment, a modulator induces a lung cancer phenotype. Generally, a plurality of assay mixtures are run in parallel with different agent concentrations to obtain a differential response to the various concentrations. Typically, one of these concentrations serves as a negative control, i.e., at zero concentration or below the level of detection.

In one aspect, a modulator will neutralize the effect of a lung cancer protein. By "neutralize" is meant that activity of a protein and the consequent effect on the cell is inhibited or blocked.

In certain embodiments, combinatorial libraries of potential modulators will be screened for an ability to bind to a lung cancer polypeptide or to modulate activity. Conventionally, new chemical entities with useful properties are generated by identifying a chemical compound (called a "lead compound") with some desirable property or activity, e.g., inhibiting activity, creating variants of the lead compound, and evaluating the property and activity of those variant compounds. Often, high throughput screening (HTS) methods are employed for such an analysis.

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In one preferred embodiment, high throughput screening methods involve providing a library containing a large number of potential therapeutic compounds (candidate compounds). Such "combinatorial chemical libraries" are then screened in one or more assays to identify those library members (particular chemical species or subclasses) that display a desired characteristic activity. The compounds thus identified can serve as conventional "lead compounds" or can themselves be used as potential or actual therapeutics.

A combinatorial chemical library is a collection of diverse chemical compounds generated by either chemical synthesis or biological synthesis by combining a number of chemical "building blocks" such as reagents. For example, a linear combinatorial chemical library, such as a polypeptide (e.g., mutein) library, is formed by combining a set of chemical building blocks called amino acids in every possible way for a given compound length (i.e., the number of amino acids in a polypeptide compound). Millions of chemical compounds can be synthesized through such combinatorial mixing of chemical building blocks (Gallop, et al. (1994) J. Med. Chem. 37(9):1233-1251).

Preparation and screening of combinatorial chemical libraries is well known to those of skill in the art. Such combinatorial chemical libraries include, but are not limited to, peptide libraries (see, e.g., U.S. Patent No. 5,010,175, Furka (1991) Pept. Prot. Res. 37:487-493, Houghton, et al. (1991) Nature, 354:84-88), peptoids (PCT Publication No WO 91/19735), encoded peptides (PCT Publication WO 93/20242), random bio-oligomers (PCT Publication WO 92/00091), benzodiazepines (U.S. Pat. No. 5,288,514), diversomers such as hydantoins, benzodiazepines and dipeptides (Hobbs, et al. (1993) Proc. Nat. Acad. Sci. USA 90:6909-6913), vinylogous polypeptides (Hagihara, et al. (1992) J. Amer. Chem. Soc. 114:6568), nonpeptidal peptidomimetics with a Beta-D-Glucose scaffolding (Hirschmann, et

al. (1992) <u>J. Amer. Chem. Soc.</u> 114:9217-9218), analogous organic syntheses of small compound libraries (Chen, et al. (1994) <u>J. Amer. Chem. Soc.</u> 116:2661), oligocarbamates (Cho, et al. (1993) <u>Science</u> 261:1303), and/or peptidyl phosphonates (Campbell, et al. (1994) <u>J. Org. Chem.</u> 59:658). See, generally, Gordon, et al. (1994) <u>J. Med. Chem.</u> 37:1385, nucleic acid libraries (see, e.g., Stratagene, Corp.), peptide nucleic acid libraries (see, e.g., U.S. Patent 5,539,083), antibody libraries (see, e.g., Vaughn, et al. (1996) <u>Nature Biotechnology</u> 14(3):309-314, and PCT/US96/10287), carbohydrate libraries (see, e.g., Liang, et al. (1996) <u>Science</u> 274:1520-1522, and U.S. Patent No. 5,593,853), and small organic molecule libraries (see, e.g., benzodiazepines, Baum (1993) C&EN, Jan 18, page 33; isoprenoids, U.S. Patent No. 5,569,588; thiazolidinones and metathiazanones, U.S. Patent No. 5,549,974; pyrrolidines, U.S. Patent Nos. 5,525,735 and 5,519,134; morpholino compounds, U.S. Patent No. 5,506,337; benzodiazepines, U.S. Patent No. 5,288,514; and the like).

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Devices for the preparation of combinatorial libraries are commercially available (see, e.g., 357 MPS, 390 MPS, Advanced Chem Tech, Louisville KY, Symphony, Rainin, Woburn, MA, 433A Applied Biosystems, Foster City, CA, 9050 Plus, Millipore, Bedford, MA).

A number of well known robotic systems have also been developed for solution phase chemistries. These systems include automated workstations like the automated synthesis apparatus developed by Takeda Chemical Industries, LTD. (Osaka, Japan) and many robotic systems utilizing robotic arms (Zymate II, Zymark Corporation, Hopkinton, Mass.; Orca, Hewlett-Packard, Palo Alto, Calif.), which mimic the manual synthetic operations performed by a chemist. The above devices, with appropriate modification, are suitable for use with the present invention. In addition, numerous combinatorial libraries are themselves commercially available (see, e.g., ComGenex, Princeton, N.J., Asinex, Moscow, Ru, Tripos, Inc., St. Louis, MO, ChemStar, Ltd, Moscow, RU, 3D Pharmaceuticals, Exton, PA, Martek Biosciences, Columbia, MD, etc.).

The assays to identify modulators are amenable to high throughput screening.

Preferred assays thus detect modulation of lung cancer gene transcription, polypeptide expression, and polypeptide activity.

High throughput assays for evaluating the presence, absence, quantification, or other properties of particular nucleic acids or protein products are well known to those of skill in the art. Similarly, binding assays and reporter gene assays are similarly well known. Thus, e.g., U.S. Patent No. 5,559,410 discloses high throughput screening methods for proteins,

U.S. Patent No. 5,585,639 discloses high throughput screening methods for nucleic acid binding (i.e., in arrays), while U.S. Patent Nos. 5,576,220 and 5,541,061 disclose high throughput methods of screening for ligand/antibody binding.

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In addition, high throughput screening systems are commercially available (see, e.g., Zymark Corp., Hopkinton, MA; Air Technical Industries, Mentor, OH; Beckman Instruments, Inc. Fullerton, CA; Precision Systems, Inc., Natick, MA, etc.). These systems typically automate procedures, including sample and reagent pipetting, liquid dispensing, timed incubations, and final readings of the microplate in detector(s) appropriate for the assay. These configurable systems provide high throughput and rapid start up as well as a high degree of flexibility and customization. The manufacturers of such systems provide detailed protocols for various high throughput systems. Thus, e.g., Zymark Corp. provides technical bulletins describing screening systems for detecting the modulation of gene transcription, ligand binding, and the like.

In one embodiment, modulators are proteins, often naturally occurring proteins or fragments of naturally occurring proteins. Thus, e.g., cellular extracts containing proteins, or random or directed digests of proteinaceous cellular extracts, may be used. In this way libraries of proteins may be made for screening in the methods of the invention. Particularly preferred in this embodiment are libraries of bacterial, fungal, viral, and mammalian proteins, with the latter being preferred, and human proteins being especially preferred. Particularly useful test compound will be directed to the class of proteins to which the target belongs, e.g., substrates for enzymes or ligands and receptors.

In a preferred embodiment, modulators are peptides of from about 5 to about 30 amino acids, with from about 5 to about 20 amino acids being preferred, and from about 7 to about 15 being particularly preferred. The peptides may be digests of naturally occurring proteins, random peptides, or "biased" random peptides. By "randomized" or grammatical equivalents herein is meant that the nucleic acid or peptide consists of essentially random sequences of nucleotides and amino acids, respectively. Since these random peptides (or nucleic acids, discussed below) are often chemically synthesized, they may incorporate a nucleotide or amino acid at any position. The synthetic process can be designed to generate randomized proteins or nucleic acids, to allow the formation of all or most of the possible combinations over the length of the sequence, thus forming a library of randomized candidate bioactive proteinaceous agents.

In one embodiment, the library is fully randomized, with no sequence preferences or constants at any position. In a preferred embodiment, the library is biased. That is, some positions within the sequence are either held constant, or are selected from a limited number of possibilities. In a preferred embodiment, the nucleotides or amino acid residues are randomized within a defined class, e.g., of hydrophobic amino acids, hydrophilic residues, sterically biased (either small or large) residues, towards the creation of nucleic acid binding domains, the creation of cysteines, for cross-linking, prolines for SH-3 domains, serines, threonines, tyrosines or histidines for phosphorylation sites, etc.

Modulators of lung cancer can also be nucleic acids, as defined above.

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As described above generally for proteins, nucleic acid modulating agents may be naturally occurring nucleic acids, random nucleic acids, or "biased" random nucleic acids. Digests of procaryotic or eucaryotic genomes may be used as is outlined above for proteins.

In a preferred embodiment, the candidate compounds are organic chemical moieties, a wide variety of which are available in the literature.

After a candidate agent has been added and the cells allowed to incubate for some period of time, the sample containing a target sequence is analyzed. If required, the target sequence is prepared using known techniques. For example, the sample may be treated to lyse the cells, using known lysis buffers, electroporation, etc., with purification and/or amplification such as PCR performed as appropriate. For example, an *in vitro* transcription with labels covalently attached to the nucleotides is performed. Generally, the nucleic acids are labeled with biotin-FITC or PE, or with cy3 or cy5.

In a preferred embodiment, the target sequence is labeled with, e.g., a fluorescent, a chemiluminescent, a chemical, or a radioactive signal, to provide a means of detecting the target sequence's specific binding to a probe. The label also can be an enzyme, such as, alkaline phosphatase or horseradish peroxidase, which when provided with an appropriate substrate produces a product that can be detected. Alternatively, the label can be a labeled compound or small molecule, such as an enzyme inhibitor, that binds but is not catalyzed or altered by the enzyme. The label also can be a moiety or compound, such as, an epitope tag or biotin which specifically binds to streptavidin. For the example of biotin, the streptavidin is labeled as described above, thereby, providing a detectable signal for the bound target sequence. Unbound labeled streptavidin is typically removed prior to analysis.

Nucleic acid assays can be direct hybridization assays or can comprise "sandwich assays", which include the use of multiple probes, as is generally outlined in U.S. Patent Nos.

5,681,702, 5,597,909, 5,545,730, 5,594,117, 5,591,584, 5,571,670, 5,580,731, 5,571,670, 5,591,584, 5,624,802, 5,635,352, 5,594,118, 5,359,100, 5,124,246 and 5,681,697, all of which are hereby incorporated by reference. In this embodiment, in general, the target nucleic acid is prepared as outlined above, and then added to the biochip comprising a plurality of nucleic acid probes, under conditions that allow the formation of a hybridization complex.

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A variety of hybridization conditions may be used in the present invention, including high, moderate and low stringency conditions as outlined above. The assays are generally run under stringency conditions which allow formation of the label probe hybridization complex only in the presence of target. Stringency can be controlled by altering a step parameter that is a thermodynamic variable, including, but not limited to, temperature, formamide concentration, salt concentration, chaotropic salt concentration, pH, organic solvent concentration, etc.

These parameters may also be used to control non-specific binding, as is generally outlined in U.S. Patent No. 5,681,697. Thus it may be desirable to perform certain steps at higher stringency conditions to reduce non-specific binding.

The reactions outlined herein may be accomplished in a variety of ways. Components of the reaction may be added simultaneously, or sequentially, in different orders, with preferred embodiments outlined below. In addition, the reaction may include a variety of other reagents. These include salts, buffers, neutral proteins, e.g., albumin, detergents, etc. which may be used to facilitate optimal hybridization and detection, and/or reduce non-specific or background interactions. Reagents that otherwise improve the efficiency of the assay, such as protease inhibitors, nuclease inhibitors, anti-microbial agents, etc., may also be used as appropriate, depending on the sample preparation methods and purity of the target.

The assay data are analyzed to determine the expression levels, and changes in expression levels as between states, of individual genes, forming a gene expression profile.

Screens are performed to identify modulators of the lung cancer phenotype. In one embodiment, screening is performed to identify modulators that can induce or suppress a particular expression profile, thus preferably generating the associated phenotype. In another embodiment, e.g., for diagnostic applications, having identified differentially expressed genes important in a particular state, screens can be performed to identify modulators that alter expression of individual genes. In an another embodiment, screening is performed to identify modulators that alter a biological function of the expression product of a differentially expressed gene. Again, having identified the importance of a gene in a particular state,

screens are performed to identify agents that bind and/or modulate the biological activity of the gene product, or evaluate genetic polymorphisms.

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Genes can be screened for those that are induced in response to a candidate agent. After identifying a modulator based upon its ability to suppress a lung cancer expression pattern leading to a normal expression pattern, or to modulate a single lung cancer gene expression profile so as to mimic the expression of the gene from normal tissue, a screen as described above can be performed to identify genes that are specifically modulated in response to the agent. Comparing expression profiles between normal tissue and agent treated lung cancer tissue reveals genes that are not expressed in normal tissue or lung cancer tissue, but are expressed in agent treated tissue. These agent-specific sequences can be identified and used by methods described herein for lung cancer genes or proteins. In particular these sequences and the proteins they encode find use in marking or identifying agent treated cells. In addition, antibodies can be raised against the agent induced proteins and used to target novel therapeutics to the treated lung cancer tissue sample.

Thus, in one embodiment, a test compound is administered to a population of lung cancer cells, that have an associated lung cancer expression profile. By "administration" or "contacting" herein is meant that the candidate agent is added to the cells in such a manner as to allow the agent to act upon the cell, whether by uptake and intracellular action, or by action at the cell surface. In some embodiments, nucleic acid encoding a proteinaceous candidate agent (i.e., a peptide) may be put into a viral construct such as an adenoviral or retroviral construct, and added to the cell, such that expression of the peptide agent is accomplished, e.g., PCT US97/01019. Regulatable gene therapy systems can also be used.

Once a test compound has been administered to the cells, the cells can be washed if desired and are allowed to incubate under preferably physiological conditions for some period of time. The cells are then harvested and a new gene expression profile is generated, as outlined herein.

Thus, e.g., lung cancer or non-malignant tissue may be screened for agents that modulate, e.g., induce or suppress a lung cancer phenotype. A change in at least one gene, preferably many, of the expression profile indicates that the agent has an effect on lung cancer activity. By defining such a signature for the lung cancer phenotype, screens for new drugs that alter the phenotype can be devised. With this approach, the drug target need not be known and need not be represented in the original expression screening platform, nor does the level of transcript for the target protein need to change.

Measure of lung cancer polypeptide activity, or of lung cancer or the lung cancer phenotype can be performed using a variety of assays. For example, the effects of the test compounds upon the function of the metastatic polypeptides can be measured by examining parameters described above. A suitable physiological change that affects activity can be used to assess the influence of a test compound on the polypeptides of this invention. When the functional consequences are determined using intact cells or animals, one can also measure a variety of effects such as, in the case of lung cancer associated with tumors, tumor growth, tumor metastasis, neovascularization, hormone release, transcriptional changes to both known and uncharacterized genetic markers (e.g., northern blots), changes in cell metabolism such as cell growth or pH changes, and changes in intracellular second messengers such as cGMP. In the assays of the invention, mammalian lung cancer polypeptide is typically used, e.g., mouse, preferably human.

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Assays to identify compounds with modulating activity can be performed *in vitro*. For example, a lung cancer polypeptide is first contacted with a potential modulator and incubated for a suitable amount of time, e.g., from 0.5 to 48 hours. In one embodiment, the lung cancer polypeptide levels are determined *in vitro* by measuring the level of protein or mRNA. The level of protein is typically measured using immunoassays such as western blotting, ELISA and the like with an antibody that selectively binds to the lung cancer polypeptide or a fragment thereof. For measurement of mRNA, amplification, e.g., using PCR, LCR, or hybridization assays, e.g., northern hybridization, RNAse protection, dot blotting, are preferred. The level of protein or mRNA is typically detected using directly or indirectly labeled detection agents, e.g., fluorescently or radioactively labeled nucleic acids, radioactively or enzymatically labeled antibodies, and the like, as described herein.

Alternatively, a reporter gene system can be devised using a lung cancer protein promoter operably linked to a reporter gene such as luciferase, green fluorescent protein, CAT, or β -gal. The reporter construct is typically transfected into a cell. After treatment with a potential modulator, the amount of reporter gene transcription, translation, or activity is measured according to standard techniques known to those of skill in the art.

In a preferred embodiment, as outlined above, screens may be done on individual genes and gene products (proteins). That is, having identified a particular differentially expressed gene as important in a particular state, screening of modulators of the expression of the gene or the gene product itself can be done. The gene products of differentially expressed

genes are sometimes referred to herein as "lung cancer proteins." The lung cancer protein may be a fragment, or alternatively, be the full length protein to a fragment shown herein.

In one embodiment, screening for modulators of expression of specific genes is performed. Typically, the expression of only one or a few genes are evaluated. In another embodiment, screens are designed to first find compounds that bind to differentially expressed proteins. These compounds are then evaluated for the ability to modulate differentially expressed activity. Moreover, once initial candidate compounds are identified, variants can be further screened to better evaluate structure activity relationships.

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In a preferred embodiment, binding assays are done. In general, purified or isolated gene product is used; that is, the gene products of one or more differentially expressed nucleic acids are made. For example, antibodies are generated to the protein gene products, and standard immunoassays are run to determine the amount of protein present. Alternatively, cells comprising the lung cancer proteins can be used in the assays.

Thus, in a preferred embodiment, the methods comprise combining a lung cancer protein and a candidate compound, and determining the binding of the compound to the lung cancer protein. Preferred embodiments utilize the human lung cancer protein, although other mammalian proteins may also be used, e.g., for the development of animal models of human disease. In some embodiments, as outlined herein, variant or derivative lung cancer proteins may be used.

Generally, in a preferred embodiment of the methods herein, the lung cancer protein or the candidate agent is non-diffusably bound to an insoluble support, preferably having isolated sample receiving areas (e.g., a microtiter plate, an array, etc.). The insoluble supports may be made of a composition to which the compositions can be bound, is readily separated from soluble material, and is otherwise compatible with the overall method of screening. The surface of such supports may be solid or porous and of a convenient shape. Examples of suitable insoluble supports include microtiter plates, arrays, membranes and beads. These are typically made of glass, plastic (e.g., polystyrene), polysaccharides, nylon or nitrocellulose, teflonTM, etc. Microtiter plates and arrays are especially convenient because a large number of assays can be carried out simultaneously, using small amounts of reagents and samples. The particular manner of binding of the composition is typically not crucial so long as it is compatible with the reagents and overall methods of the invention, maintains the activity of the composition, and is nondiffusable. Preferred methods of binding include the use of antibodies (which do not sterically block either the ligand binding site or activation

sequence when the protein is bound to the support), direct binding to "sticky" or ionic supports, chemical crosslinking, the synthesis of the protein or agent on the surface, etc. Following binding of the protein or agent, excess unbound material is removed by washing. The sample receiving areas may then be blocked through incubation with bovine serum albumin (BSA), casein or other innocuous protein or other moiety.

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In a preferred embodiment, the lung cancer protein is bound to the support, and a test compound is added to the assay. Alternatively, the candidate agent is bound to the support and the lung cancer protein is added. Novel binding agents include specific antibodies, non-natural binding agents identified in screens of chemical libraries, peptide analogs, etc. Of particular interest are screening assays for agents that have a low toxicity for human cells. A wide variety of assays may be used for this purpose, including labeled *in vitro* protein-protein binding assays, electrophoretic mobility shift assays, immunoassays for protein binding, functional assays (phosphorylation assays, etc.) and the like.

The determination of the binding of the test modulating compound to the lung cancer protein may be done in a number of ways. In a preferred embodiment, the compound is labeled, and binding determined directly, e.g., by attaching all or a portion of the lung cancer protein to a solid support, adding a labeled candidate agent (e.g., a fluorescent label), washing off excess reagent, and determining whether the label is present on the solid support. Various blocking and washing steps may be utilized as appropriate.

In some embodiments, only one of the components is labeled, e.g., the proteins (or proteinaceous candidate compounds) can be labeled. Alternatively, more than one component can be labeled with different labels, e.g., ¹²⁵I for the proteins and a fluorophor for the compound. Proximity reagents, e.g., quenching or energy transfer reagents are also useful.

In one embodiment, the binding of the test compound is determined by competitive binding assay. The competitor may be a binding moiety known to bind to the target molecule (i.e., a lung cancer protein), such as an antibody, peptide, binding partner, ligand, etc. Under certain circumstances, there may be competitive binding between the compound and the binding moiety, with the binding moiety displacing the compound. In one embodiment, the test compound is labeled. Either the compound, or the competitor, or both, is added first to the protein for a time sufficient to allow binding, if present. Incubations may be performed at a temperature which facilitates optimal activity, typically between 4 and 40° C. Incubation periods are typically optimized, e.g., to facilitate rapid high throughput screening. Typically

between 0.1 and 1 hour will be sufficient. Excess reagent is generally removed or washed away. The second component is then added, and the presence or absence of the labeled component is followed, to indicate binding.

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In a preferred embodiment, the competitor is added first, followed by a test compound. Displacement of the competitor is an indication that the test compound is binding to the lung cancer protein and thus is capable of binding to, and potentially modulating, the activity of the lung cancer protein. In this embodiment, either component can be labeled. Thus, e.g., if the competitor is labeled, the presence of label in the wash solution indicates displacement by the agent. Alternatively, if the test compound is labeled, the presence of the label on the support indicates displacement.

In an alternative embodiment, the test compound is added first, with incubation and washing, followed by the competitor. The absence of binding by the competitor may indicate that the test compound is bound to the lung cancer protein with a higher affinity. Thus, if the test compound is labeled, the presence of the label on the support, coupled with a lack of competitor binding, may indicate that the test compound is capable of binding to the lung cancer protein.

In a preferred embodiment, the methods comprise differential screening to identity agents that are capable of modulating the activity of the lung cancer proteins. In one embodiment, the methods comprise combining a lung cancer protein and a competitor in a first sample. A second sample comprises a test compound, a lung cancer protein, and a competitor. The binding of the competitor is determined for both samples, and a change, or difference in binding between the two samples indicates the presence of an agent capable of binding to the lung cancer protein and potentially modulating its activity. That is, if the binding of the competitor is different in the second sample relative to the first sample, the agent is capable of binding to the lung cancer protein.

Alternatively, differential screening is used to identify drug candidates that bind to the native lung cancer protein, but cannot bind to modified lung cancer proteins. The structure of the lung cancer protein may be modeled, and used in rational drug design to synthesize agents that interact with that site. Drug candidates that affect the activity of a lung cancer protein are also identified by screening drugs for the ability to either enhance or reduce the activity of the protein.

Positive controls and negative controls may be used in the assays. Preferably control and test samples are performed in at least triplicate to obtain statistically significant results.

Incubation of all samples is for a time sufficient for the binding of the agent to the protein. Following incubation, samples are washed free of non-specifically bound material and the amount of bound, generally labeled agent determined. For example, where a radiolabel is employed, the samples may be counted in a scintillation counter to determine the amount of bound compound.

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A variety of other reagents may be included in the screening assays. These include reagents like salts, neutral proteins, e.g., albumin, detergents, etc. which may be used to facilitate optimal protein-protein binding and/or reduce non-specific or background interactions. Also reagents that otherwise improve the efficiency of the assay, such as protease inhibitors, nuclease inhibitors, anti-microbial agents, etc., may be used. The mixture of components may be added in an order that provides for the requisite binding.

In a preferred embodiment, the invention provides methods for screening for a compound capable of modulating the activity of a lung cancer protein. The methods comprise adding a test compound, as defined above, to a cell comprising lung cancer proteins. Preferred cell types include almost any cell. The cells contain a recombinant nucleic acid that encodes a lung cancer protein. In a preferred embodiment, a library of candidate agents are tested on a plurality of cells.

In one aspect, the assays are evaluated in the presence or absence or previous or subsequent exposure of physiological signals, e.g., hormones, antibodies, peptides, antigens, cytokines, growth factors, action potentials, pharmacological agents including chemotherapeutics, radiation, carcinogenics, or other cells (e.g., cell-cell contacts). In another example, the determinations are determined at different stages of the cell cycle process.

In this way, compounds that modulate lung cancer agents are identified. Compounds with pharmacological activity are able to enhance or interfere with the activity of the lung cancer protein. Once identified, similar structures are evaluated to identify critical structural feature of the compound.

In one embodiment, a method of inhibiting lung cancer cell division is provided. The method comprises administration of a lung cancer inhibitor. In another embodiment, a method of inhibiting lung cancer is provided. The method may comprise administration of a lung cancer inhibitor. In a further embodiment, methods of treating cells or individuals with lung cancer are provided, e.g., comprising administration of a lung cancer inhibitor.

In one embodiment, a lung cancer inhibitor is an antibody as discussed above. In another embodiment, the lung cancer inhibitor is an antisense molecule.

A variety of cell growth, proliferation, viability, and metastasis assays are known to those of skill in the art, as described below.

Soft agar growth or colony formation in suspension

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Normal cells require a solid substrate to attach and grow. When the cells are transformed, they lose this phenotype and grow detached from the substrate. For example, transformed cells can grow in stirred suspension culture or suspended in semi-solid media, such as semi-solid or soft agar. The transformed cells, when transfected with tumor suppressor genes, regenerate normal phenotype and require a solid substrate to attach and grow. Soft agar growth or colony formation in suspension assays can be used to identify modulators of lung cancer sequences, which when expressed in host cells, inhibit abnormal cellular proliferation and transformation. A therapeutic compound would reduce or eliminate the host cells' ability to grow in stirred suspension culture or suspended in semi-solid media, such as semi-solid or soft.

Techniques for soft agar growth or colony formation in suspension assays are described in Freshney (1994) <u>Culture of Animal Cells a Manual of Basic Technique</u> (3rd ed.), herein incorporated by reference. See also, the methods section of Garkavtsev, et al. (1996), *supra*, herein incorporated by reference.

20 Contact inhibition and density limitation of growth

Normal cells typically grow in a flat and organized pattern in a petri dish until they touch other cells. When the cells touch one another, they are contact inhibited and stop growing. When cells are transformed, however, the cells are not contact inhibited and continue to grow to high densities in disorganized foci. Thus, the transformed cells grow to a higher saturation density than normal cells. This can be detected morphologically by the formation of a disoriented monolayer of cells or rounded cells in foci within the regular pattern of normal surrounding cells. Alternatively, labeling index with (³H)-thymidine at saturation density can be used to measure density limitation of growth. See Freshney (1994), supra. The transformed cells, when transfected with tumor suppressor genes, regenerate a normal phenotype and become contact inhibited and would grow to a lower density.

In this assay, labeling index with (³H)-thymidine at saturation density is a preferred method of measuring density limitation of growth. Transformed host cells are transfected with a lung cancer-associated sequence and are grown for 24 hours at saturation density in

non-limiting medium conditions. The percentage of cells labeling with (³H)-thymidine is determined autoradiographically. See, Freshney (1994), *supra*.

Growth factor or serum dependence

Transformed cells typically have a lower serum dependence than their normal counterparts (see, e.g., Temin (1966) <u>J. Natl. Cancer Insti.</u> 37:167-175; Eagle, et al. (1970) <u>J. Exp. Med.</u> 131:836-879); Freshney, *supra*. This is in part due to release of various growth factors by the transformed cells. Growth factor or serum dependence of transformed host cells can be compared with that of control.

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Tumor specific markers levels

Tumor cells release an increased amount of certain factors (hereinafter "tumor specific markers") than their normal counterparts. For example, plasminogen activator (PA) is released from human glioma at a higher level than from normal brain cells (see, e.g., Gullino, "Angiogenesis, tumor vascularization, and potential interference with tumor growth" in Mihich (ed. 1985) Biological Responses in Cancer, pp. 178-184). Similarly, Tumor angiogenesis factor (TAF) is released at a higher level in tumor cells than their normal counterparts. See, e.g., Folkman (1992) "Angiogenesis and Cancer" in Sem Cancer Biol.).

Various techniques which measure the release of these factors are described in Freshney (1994), *supra*. Also, see, Unkeless, et al. (1974) <u>J. Biol. Chem.</u> 249:4295-4305; Strickland and Beers (1976) <u>J. Biol. Chem.</u> 251:5694-5702; Whur, et al. (1980) <u>Br. J. Cancer</u> 42:305-312; Gullino, "Angiogenesis, tumor vascularization, and potential interference with tumor growth" in Mihich (ed. 1985) <u>Biological Responses in Cancer</u>, pp. 178-184; Freshney <u>Anticancer Res.</u> 5:111-130 (1985).

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Invasiveness into Matrigel

The degree of invasiveness into Matrigel or some other extracellular matrix constituent can be used as an assay to identify compounds that modulate lung cancer-associated sequences. Tumor cells exhibit a good correlation between malignancy and invasiveness of cells into Matrigel or some other extracellular matrix constituent. In this assay, tumorigenic cells are typically used as host cells. Expression of a tumor suppressor gene in these host cells would decrease invasiveness of the host cells.

Techniques described in Freshney (1994), *supra*, can be used. Briefly, the level of invasion of host cells can be measured by using filters coated with Matrigel or some other extracellular matrix constituent. Penetration into the gel, or through to the distal side of the filter, is rated as invasiveness, and rated histologically by number of cells and distance moved, or by prelabeling the cells with ¹²⁵I and counting the radioactivity on the distal side of the filter or bottom of the dish. See, e.g., Freshney (1984), *supra*.

Tumor growth in vivo

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Effects of lung cancer-associated sequences on cell growth can be tested in transgenic or immune-suppressed mice. Knock-out transgenic mice can be made, in which the lung cancer gene is disrupted or in which a lung cancer gene is inserted. Knock-out transgenic mice can be made by insertion of a marker gene or other heterologous gene into the endogenous lung cancer gene site in the mouse genome via homologous recombination. Such mice can also be made by substituting the endogenous lung cancer gene with a mutated version of the lung cancer gene, or by mutating the endogenous lung cancer gene, e.g., by exposure to carcinogens.

A DNA construct is introduced into the nuclei of embryonic stem cells. Cells containing the newly engineered genetic lesion are injected into a host mouse embryo, which is re-implanted into a recipient female. Some of these embryos develop into chimeric mice that possess germ cells partially derived from the mutant cell line. Therefore, by breeding the chimeric mice it is possible to obtain a new line of mice containing the introduced genetic lesion (see, e.g., Capecchi, et al. (1989) Science 244:1288). Chimeric targeted mice can be derived according to Hogan, et al. (1988) Manipulating the Mouse Embryo: A Laboratory Manual, Cold Spring Harbor Laboratory and Robertson (ed. 1987) Teratocarcinomas and Embryonic Stem Cells: A Practical Approach, , IRL Press, Washington, D.C.

Alternatively, various immune-suppressed or immune-deficient host animals can be used. For example, genetically athymic "nude" mouse (see, e.g., Giovanella, et al. (1974) <u>J. Natl. Cancer Inst.</u> 52:921), a SCID mouse, a thymectomized mouse, or an irradiated mouse (see, e.g., Bradley, et al. (1978) <u>Br. J. Cancer</u> 38:263; Selby, et al. (1980) <u>Br. J. Cancer</u> 41:52) can be used as a host. Transplantable tumor cells (typically about 10⁶ cells) injected into isogenic hosts will produce invasive tumors in a high proportions of cases, while normal cells of similar origin will not. In hosts which developed invasive tumors, cells expressing a lung cancer-associated sequences are injected subcutaneously. After a suitable length of time,

preferably 4-8 weeks, tumor growth is measured (e.g., by volume or by its two largest dimensions) and compared to the control. Tumors that have statistically significant reduction (using, e.g., Student's T test) are said to have inhibited growth.

5 Polynucleotide modulators of lung cancer

Antisense and RNAi Polynucleotides

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In certain embodiments, the activity of a lung cancer-associated protein is downregulated, or entirely inhibited, by the use of antisense or an inhibitory polynucleotide, i.e., a nucleic acid complementary to, and which can preferably hybridize specifically to, a coding mRNA nucleic acid sequence, e.g., a lung cancer protein mRNA, or a subsequence thereof. Binding of the antisense polynucleotide to the mRNA reduces the translation and/or stability of the mRNA.

In the context of this invention, antisense polynucleotides can comprise naturally-occurring nucleotides, or synthetic species formed from naturally-occurring subunits or their close homologs. Antisense polynucleotides may also have altered sugar moieties or intersugar linkages. Exemplary among these are the phosphorothioate and other sulfur containing species which are known for use in the art. Analogs are comprehended by this invention so long as they function effectively to hybridize with the lung cancer protein mRNA. See, e.g., Isis Pharmaceuticals, Carlsbad, CA; Sequitor, Inc., Natick, MA.

Such antisense polynucleotides can readily be synthesized using recombinant means, or can be synthesized *in vitro*. Equipment for such synthesis is sold by several vendors, including Applied Biosystems. The preparation of other oligonucleotides such as phosphorothioates and alkylated derivatives is also well known to those of skill in the art.

Antisense molecules as used herein include antisense or sense oligonucleotides. Sense oligonucleotides can, e.g., be employed to block transcription by binding to the antisense strand. The antisense and sense oligonucleotide comprise a single-stranded nucleic acid sequence (either RNA or DNA) capable of binding to target mRNA (sense) or DNA (antisense) sequences for lung cancer molecules. A preferred antisense molecule is for a lung cancer sequence in the tables, or for a ligand or activator thereof. Antisense or sense oligonucleotides, according to the present invention, comprise a fragment generally at least about 14 nucleotides, preferably from about 14 to 30 nucleotides. The ability to derive an antisense or a sense oligonucleotide, based upon a cDNA sequence encoding a given protein

WO 02/086443
PCT/US02/12476
is described in, e.g., Stein and Cohen (1988) Cancer Res. 48:2659 and van der Krol, et al.
(1988) BioTechniques 6:958).

RNA interference is a mechanism to suppress gene expression in a sequence specific manner. See, e.g., Brumelkamp, et al. (2002) Sciencexpress (21March2002); Sharp (1999) Genes Dev. 13:139-141; and Cathew (2001) Curr. Op. Cell Biol. 13:244-248. In mammalian cells, short, e.g., 21 nt, double stranded small interfering RNAs (siRNA) have been shown to be effective at inducing an RNAi response. See, e.g., Elbashir, et al. (2001) Nature 411:494-498. The mechanism may be used to downregulate expression levels of identified genes, e.g., treatment of or validation of relevance to disease.

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Ribozymes

In addition to antisense polynucleotides, ribozymes can be used to target and inhibit transcription of lung cancer-associated nucleotide sequences. A ribozyme is an RNA molecule that catalytically cleaves other RNA molecules. Different kinds of ribozymes have been described, including group I ribozymes, hammerhead ribozymes, hairpin ribozymes, RNase P, and axhead ribozymes (see, e.g., Castanotto, et al. (1994) Adv. in Pharmacology 25: 289-317 for a general review of the properties of different ribozymes).

The general features of hairpin ribozymes are described, e.g., in Hampel, et al. (1990) Nucl. Acids Res. 18:299-304; European Patent Publication No. 0 360 257; U.S. Patent No. 5,254,678. Methods of preparing are well known to those of skill in the art (see, e.g., WO 94/26877; Ojwang, et al. (1993) Proc. Natl. Acad. Sci. USA 90:6340-6344; Yamada, et al. (1994) Human Gene Therapy 1:39-45; Leavitt, et al. (1995) Proc. Natl. Acad. Sci. USA 92:699-703; Leavitt, et al. (1994) Human Gene Therapy 5:1151-120; and Yamada, et al. (1994) Virology 205: 121-126).

Polynucleotide modulators of lung cancer may be introduced into a cell containing the target nucleotide sequence by formation of a conjugate with a ligand binding molecule, as described in WO 91/04753. Suitable ligand binding molecules include, but are not limited to, cell surface receptors, growth factors, other cytokines, or other ligands that bind to cell surface receptors. Preferably, conjugation of the ligand binding molecule does not substantially interfere with the ability of the ligand binding molecule to bind to its corresponding molecule or receptor, or block entry of the sense or antisense oligonucleotide or its conjugated version into the cell. Alternatively, a polynucleotide modulator of lung cancer may be introduced into a cell containing the target nucleic acid sequence, e.g., by

formation of an polynucleotide-lipid complex, as described in WO 90/10448. It is understood that the use of antisense molecules or knock out and knock in models may also be used in screening assays as discussed above, in addition to methods of treatment.

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Thus, in one embodiment, methods of modulating lung cancer in cells or organisms are provided. In one embodiment, the methods comprise administering to a cell an anti-lung cancer antibody that reduces or eliminates the biological activity of an endogenous lung cancer protein. Alternatively, the methods comprise administering to a cell or organism a recombinant nucleic acid encoding a lung cancer protein. This may be accomplished in any number of ways. In a preferred embodiment, e.g., when the lung cancer sequence is down-regulated in lung cancer, such state may be reversed by increasing the amount of lung cancer gene product in the cell. This can be accomplished, e.g., by overexpressing the endogenous lung cancer gene or administering a gene encoding the lung cancer sequence, using known gene-therapy techniques. In a preferred embodiment, the gene therapy techniques include the incorporation of the exogenous gene using enhanced homologous recombination (EHR), e.g., as described in PCT/US93/03868, hereby incorporated by reference in its entirety. Alternatively, e.g., when the lung cancer sequence is up-regulated in lung cancer, the activity of the endogenous lung cancer gene is decreased, e.g., by the administration of a lung cancer antisense or RNAi nucleic acid.

In one embodiment, the lung cancer proteins of the present invention may be used to generate polyclonal and monoclonal antibodies to lung cancer proteins. Similarly, the lung cancer proteins can be coupled, using standard technology, to affinity chromatography columns. These columns may then be used to purify lung cancer antibodies useful for production, diagnostic, or therapeutic purposes. In a preferred embodiment, the antibodies are generated to epitopes unique to a lung cancer protein; that is, the antibodies show little or no cross-reactivity to other proteins. The lung cancer antibodies may be coupled to standard affinity chromatography columns and used to purify lung cancer proteins. The antibodies may also be used as blocking polypeptides, as outlined above, since they will specifically bind to the lung cancer protein.

Methods of identifying variant lung cancer-associated sequences

Without being bound by theory, expression of various lung cancer sequences is correlated with lung cancer. Accordingly, disorders based on mutant or variant lung cancer genes may be determined. In one embodiment, the invention provides methods for

identifying cells containing variant lung cancer genes, e.g., determining all or part of the sequence of at least one endogenous lung cancer genes in a cell. In a preferred embodiment, the invention provides methods of identifying the lung cancer genotype of an individual, e.g., determining all or part of the sequence of at least one lung cancer gene of the individual. This is generally done in at least one tissue of the individual, and may include the evaluation of a number of tissues or different samples of the same tissue. The method may include comparing the sequence of the sequenced lung cancer gene to a known lung cancer gene, i.e.,

The sequence of all or part of the lung cancer gene can then be compared to the sequence of a known lung cancer gene to determine if any differences exist. This can be done using known homology programs, such as Bestfit, etc. In a preferred embodiment, the presence of a difference in the sequence between the lung cancer gene of the patient and the known lung cancer gene correlates with a disease state or a propensity for a disease state, as outlined herein.

In a preferred embodiment, the lung cancer genes are used as probes to determine the number of copies of the lung cancer gene in the genome.

In another preferred embodiment, the lung cancer genes are used as probes to determine the chromosomal localization of the lung cancer genes. Information such as chromosomal localization finds use in providing a diagnosis or prognosis in particular when chromosomal abnormalities such as translocations, and the like are identified in the lung cancer gene locus.

Administration of pharmaceutical and vaccine compositions

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a wild-type gene.

In one embodiment, a therapeutically effective dose of a lung cancer protein or modulator thereof, is administered to a patient. By "therapeutically effective dose" herein is meant a dose that produces effects for which it is administered. The exact dose will depend on the purpose of the treatment, and will be ascertainable by one skilled in the art using known techniques (e.g., Ansel, et al. (1992) Pharmaceutical Dosage Forms and Drug Delivery; Lieberman, Pharmaceutical Dosage Forms (vols. 1-3), Dekker, ISBN 0824770846, 082476918X, 0824712692, 0824716981; Lloyd (1999) The Art. Science and Technology of Pharmaceutical Compounding; and Pickar (1999) Dosage Calculations). Adjustments for lung cancer degradation, systemic versus localized delivery, and rate of new protease synthesis, as well as the age, body weight, general health, sex, diet, time of administration,

drug interaction and the severity of the condition may be necessary, and will be ascertainable with routine experimentation by those skilled in the art.

A "patient" for the purposes of the present invention includes both humans and other animals, particularly mammals. Thus the methods are applicable to both human therapy and veterinary applications. In the preferred embodiment the patient is a mammal, preferably a primate, and in the most preferred embodiment the patient is human.

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The administration of the lung cancer proteins and modulators thereof of the present invention can be done in a variety of ways, including, but not limited to, orally, subcutaneously, intravenously, intranasally, transdermally, intraperitoneally, intramuscularly, intrapulmonary, vaginally, rectally, or intraocularly. In some instances, e.g., in the treatment of wounds and inflammation, the lung cancer proteins and modulators may be directly applied as a solution or spray.

The pharmaceutical compositions of the present invention comprise a lung cancer protein in a form suitable for administration to a patient. In the preferred embodiment, the pharmaceutical compositions are in a water soluble form, such as being present as pharmaceutically acceptable salts, which is meant to include both acid and base addition salts. "Pharmaceutically acceptable acid addition salt" refers to those salts that retain the biological effectiveness of the free bases and that are not biologically or otherwise undesirable, formed with inorganic acids such as hydrochloric acid, hydrobromic acid, sulfuric acid, nitric acid, phosphoric acid and the like, and organic acids such as acetic acid, propionic acid, glycolic acid, pyruvic acid, oxalic acid, maleic acid, malonic acid, succinic acid, fumaric acid, tartaric acid, citric acid, benzoic acid, cinnamic acid, mandelic acid, methanesulfonic acid, ethanesulfonic acid, p-toluenesulfonic acid, salicylic acid and the like. "Pharmaceutically acceptable base addition salts" include those derived from inorganic bases such as sodium, potassium, lithium, ammonium, calcium, magnesium, iron, zinc, copper, manganese, aluminum salts and the like. Particularly preferred are the ammonium, potassium, sodium, calcium, and magnesium salts. Salts derived from pharmaceutically acceptable organic non-toxic bases include salts of primary, secondary, and tertiary amines, substituted amines including naturally occurring substituted amines, cyclic amines and basic ion exchange resins, such as isopropylamine, trimethylamine, diethylamine, triethylamine, tripropylamine, and ethanolamine.

The pharmaceutical compositions may also include one or more of the following: carrier proteins such as serum albumin; buffers; fillers such as microcrystalline cellulose,

lactose, corn and other starches; binding agents; sweeteners and other flavoring agents; coloring agents; and polyethylene glycol.

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The pharmaceutical compositions can be administered in a variety of unit dosage forms depending upon the method of administration. For example, unit dosage forms suitable for oral administration include, but are not limited to, powder, tablets, pills, capsules and lozenges. It is recognized that lung cancer protein modulators (e.g., antibodies, antisense constructs, ribozymes, small organic molecules, etc.) when administered orally, should be protected from digestion. This is typically accomplished either by complexing the molecule(s) with a composition to render it resistant to acidic and enzymatic hydrolysis, or by packaging the molecule(s) in an appropriately resistant carrier, such as a liposome or a protection barrier. Means of protecting agents from digestion are well known in the art.

The compositions for administration will commonly comprise a lung cancer protein modulator dissolved in a pharmaceutically acceptable carrier, preferably an aqueous carrier. A variety of aqueous carriers can be used, e.g., buffered saline and the like. These solutions are sterile and generally free of undesirable matter. These compositions may be sterilized by conventional, well known sterilization techniques. The compositions may contain pharmaceutically acceptable auxiliary substances as required to approximate physiological conditions such as pH adjusting and buffering agents, toxicity adjusting agents and the like, e.g., sodium acetate, sodium chloride, potassium chloride, calcium chloride, sodium lactate and the like. The concentration of active agent in these formulations can vary widely, and will be selected primarily based on fluid volumes, viscosities, body weight and the like in accordance with the particular mode of administration selected and the patient's needs (e.g., Remington's Pharmaceutical Science (15th ed., 1980) and Hardman, et al. (eds. 1996) Goodman and Gilman: The Pharmacologial Basis of Therapeutics).

Thus, a typical pharmaceutical composition for intravenous administration would be about 0.1 to 10 mg per patient per day. Dosages from 0.1 up to about 100 mg per patient per day may be used, particularly when the drug is administered to a secluded site and not into the blood stream, such as into a body cavity or into a lumen of an organ. Substantially higher dosages are possible in topical administration. Actual methods for preparing parenterally administrable compositions will be known or apparent to those skilled in the art, e.g., Remington's Pharmaceutical Science and Goodman and Gilman, The Pharmacologial Basis of Therapeutics, supra.

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The compositions containing modulators of lung cancer proteins can be administered for therapeutic or prophylactic treatments. In therapeutic applications, compositions are administered to a patient suffering from a disease (e.g., a cancer) in an amount sufficient to cure or at least partially arrest the disease and its complications. An amount adequate to accomplish this is defined as a "therapeutically effective dose." Amounts effective for this use will depend upon the severity of the disease and the general state of the patient's health. Single or multiple administrations of the compositions may be administered depending on the dosage and frequency as required and tolerated by the patient. In any event, the composition should provide a sufficient quantity of the agents of this invention to effectively treat the patient. An amount of modulator that is capable of preventing or slowing the development of cancer in a mammal is referred to as a "prophylactically effective dose." The particular dose required for a prophylactic treatment will depend upon the medical condition and history of the mammal, the particular cancer being prevented, as well as other factors such as age, weight, gender, administration route, efficiency, etc. Such prophylactic treatments may be used, e.g., in a mammal who has previously had cancer to prevent a recurrence of the cancer, or in a mammal who is suspected of having a significant likelihood of developing cancer based, at least in part, upon gene expression profiles. Vaccine strategies may be used, in either a DNA vaccine form, or protein vaccine.

It will be appreciated that the present lung cancer protein-modulating compounds can be administered alone or in combination with additional lung cancer modulating compounds or with other therapeutic agent, e.g., other anti-cancer agents or treatments.

In numerous embodiments, one or more nucleic acids, e.g., polynucleotides comprising nucleic acid sequences set forth in the tables, such as antisense or RNAi polynucleotides or ribozymes, will be introduced into cells, *in vitro* or *in vivo*. The present invention provides methods, reagents, vectors, and cells useful for expression of lung cancer-associated polypeptides and nucleic acids using *in vitro* (cell-free), *ex vivo*, or *in vivo* (cell or organism-based) recombinant expression systems.

The particular procedure used to introduce the nucleic acids into a host cell for expression of a protein or nucleic acid is application specific. Many procedures for introducing foreign nucleotide sequences into host cells may be used. These include the use of calcium phosphate transfection, spheroplasts, electroporation, liposomes, microinjection, plasma vectors, viral vectors and other well known methods for introducing cloned genomic DNA, cDNA, synthetic DNA or other foreign genetic material into a host cell (see, e.g.,

Berger and Kimmel, <u>Guide to Molecular Cloning Techniques</u>, <u>Methods in Enzymology</u> volume 152 (Berger), Ausubel, et al. (eds. 1999) <u>Current Protocols</u> (supplemented through 1999), and Sambrook, et al. (1989) <u>Molecular Cloning - A Laboratory Manual</u> (2nd ed., Vol. 1-3).

In a preferred embodiment, lung cancer proteins and modulators are administered as therapeutic agents, and can be formulated as outlined above. Similarly, lung cancer genes (including both the full-length sequence, partial sequences, or regulatory sequences of the lung cancer coding regions) can be administered in a gene therapy application. These lung cancer genes can include antisense or inhibitory applications, e.g., as inhibitory RNA or gene therapy (e.g., for incorporation into the genome) or as antisense compositions.

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Lung cancer polypeptides and polynucleotides can also be administered as vaccine compositions to stimulate HTL, CTL, and antibody responses.. Such vaccine compositions can include, e.g., lipidated peptides (see, e.g., Vitiello, et al. (1995) J. Clin. Invest. 95:341), peptide compositions encapsulated in poly(DL-lactide-co-glycolide) ("PLG") microspheres (see, e.g., Eldridge, et al. (1991) Molec. Immunol. 28:287-294; Alonso, et al. (1994) Vaccine 12:299-306; Jones, et al. (1995) Vaccine 13:675-681), peptide compositions contained in immune stimulating complexes (ISCOMS) (see, e.g., Takahashi, et al. (1990) Nature 344:873-875; Hu, et al. (1998) Clin Exp Immunol. 113:235-243), multiple antigen peptide systems (MAPs) (see, e.g., Tam (1988) Proc. Natl. Acad. Sci. U.S.A. 85:5409-5413; Tam (1996) J. Immunol. Methods 196:17-32), peptides formulated as multivalent peptides; peptides for use in ballistic delivery systems, typically crystallized peptides, viral delivery vectors (Perkus, et al., p. 379 In: Kaufmann (ed. 1996) Concepts in vaccine development; Chakrabarti, et al. (1986) Nature 320:535; Hu, et al. (1986) Nature 320:537; Kieny, et al. (1986) AIDS Bio/Technology 4:790; Top, et al. (1971) J. Infect. Dis. 124:148; Chanda, et al. (1990) Virology 175:535), particles of viral or synthetic origin (see, e.g., Kofler, et al. (1996) J. Immunol. Methods 192:25; Eldridge, et al. (1993) Sem. Hematol. 30:16; Falo, et al. (1995) Nature Med. 7:649), adjuvants (Warren, et al. (1986) Annu. Rev. Immunol. 4:369; Gupta, et al. (1993) Vaccine 11:293), liposomes (Reddy, et al. (1992) J. Immunol. 148:1585; Rock (1996) Immunol. Today 17:131), or, naked or particle absorbed cDNA (Ulmer, et al. (1993) Science 259:1745; Robinson, et al. (1993) Vaccine 11:957; Shiver, et al., p. 423 In: Kaufmann (ed. 1996) Concepts in vaccine development; Cease and Berzofsky (1994) Annu. Rev. Immunol. 12:923 and Eldridge, et al. (1993) Sem. Hematol. 30:16). Toxin-targeted

delivery technologies, also known as receptor mediated targeting, such as those of Avant Immunotherapeutics, Inc. (Needham, Massachusetts) may also be used.

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Vaccine compositions often include adjuvants. Many adjuvants contain a substance designed to protect the antigen from rapid catabolism, such as aluminum hydroxide or mineral oil, and a stimulator of immune responses, such as lipid A, *Bortadella pertussis* or *Mycobacterium tuberculosis* derived proteins. Certain adjuvants are commercially available as, e.g., Freund's Incomplete Adjuvant and Complete Adjuvant (Difco Laboratories, Detroit, MI); Merck Adjuvant 65 (Merck and Company, Inc., Rahway, NJ); AS-2 (SmithKline Beecham, Philadelphia, PA); aluminum salts such as aluminum hydroxide gel (alum) or aluminum phosphate; salts of calcium, iron or zinc; an insoluble suspension of acylated tyrosine; acylated sugars; cationically or anionically derivatized polysaccharides; polyphosphazenes; biodegradable microspheres; monophosphoryl lipid A and quil A. Cytokines, such as GM-CSF, interleukin-2, -7, -12, and other like growth factors, may also be used as adjuvants.

Vaccines can be administered as nucleic acid compositions wherein DNA or RNA encoding one or more of the polypeptides, or a fragment thereof, is administered to a patient. This approach is described, for instance, in Wolff, et. al. (1990) Science 247:1465 as well as U.S. Patent Nos. 5,580,859; 5,589,466; 5,804,566; 5,739,118; 5,736,524; 5,679,647; WO 98/04720; and in more detail below. Examples of DNA-based delivery technologies include "naked DNA", facilitated (bupivicaine, polymers, peptide-mediated) delivery, cationic lipid complexes, and particle-mediated ("gene gun") or pressure-mediated delivery (see, e.g., U.S. Patent No. 5,922,687).

For therapeutic or prophylactic immunization purposes, the peptides of the invention can be expressed by viral or bacterial vectors. Examples of expression vectors include attenuated viral hosts, such as vaccinia or fowlpox. This approach involves the use of vaccinia virus, e.g., as a vector to express nucleotide sequences that encode lung cancer polypeptides or polypeptide fragments. Upon introduction into a host, the recombinant vaccinia virus expresses the immunogenic peptide, and thereby elicits an immune response. Vaccinia vectors and methods useful in immunization protocols are described in, e.g., U.S. Patent No. 4,722,848. Another vector is BCG (Bacille Calmette Guerin). BCG vectors are described in Stover, et al. (1991) Nature 351:456-460. A wide variety of other vectors useful for therapeutic administration or immunization e.g., adeno and adeno-associated virus vectors, retroviral vectors, Salmonella typhi vectors, detoxified anthrax toxin vectors, and the

like, will be apparent to those skilled in the art from the description herein (see, e.g., Shata, et al. (2000) Mol Med Today 6:66-71; Shedlock, et al. (2000) J. Leukoc. Biol. 68:793-806; Hipp, et al. (2000) In Vivo 14:571-85).

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Methods for the use of genes as DNA vaccines are well known, and include placing a lung cancer gene or portion of a lung cancer gene under the control of a regulatable promoter or a tissue-specific promoter for expression in a lung cancer patient. The lung cancer gene used for DNA vaccines can encode full-length lung cancer proteins, but more preferably encodes portions of the lung cancer proteins including peptides derived from the lung cancer protein. In one embodiment, a patient is immunized with a DNA vaccine comprising a plurality of nucleotide sequences derived from a lung cancer gene. For example, lung cancer-associated genes or sequence encoding subfragments of a lung cancer protein are introduced into expression vectors and tested for their immunogenicity in the context of Class I MHC and an ability to generate cytotoxic T cell responses. This procedure provides for production of cytotoxic T cell responses against cells which present antigen, including intracellular epitopes.

In a preferred embodiment, DNA vaccines include a gene encoding an adjuvant molecule with the DNA vaccine. Such adjuvant molecules include cytokines that increase the immunogenic response to the lung cancer polypeptide encoded by the DNA vaccine. Additional or alternative adjuvants are available.

In another preferred embodiment lung cancer genes find use in generating animal models of lung cancer. When the lung cancer gene identified is repressed or diminished in metastatic tissue, gene therapy technology, e.g., wherein antisense or inhibitory RNA directed to the lung cancer gene will also diminish or repress expression of the gene. Animal models of lung cancer find use in screening for modulators of a lung cancer-associated sequence or modulators of lung cancer. Similarly, transgenic animal technology including gene knockout technology, e.g., as a result of homologous recombination with an appropriate gene targeting vector, will result in the absence or increased expression of the lung cancer protein. When desired, tissue-specific expression or knockout of the lung cancer protein may be necessary.

It is also possible that the lung cancer protein is overexpressed in lung cancer. As such, transgenic animals can be generated that overexpress the lung cancer protein.

Depending on the desired expression level, promoters of various strengths can be employed to express the transgene. Also, the number of copies of the integrated transgene can be determined and compared for a determination of the expression level of the transgene.

Animals generated by such methods will find use as animal models of lung cancer and are additionally useful in screening for modulators to treat lung cancer.

Kits for Use in Diagnostic and/or Prognostic Applications

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For use in diagnostic, research, and therapeutic applications suggested above, kits are also provided by the invention. In diagnostic and research applications such kits may include at least one of the following: assay reagents, buffers, lung cancer-specific nucleic acids or antibodies, hybridization probes and/or primers, antisense polynucleotides, ribozymes, RNAi, dominant negative lung cancer polypeptides or polynucleotides, small molecule inhibitors of lung cancer-associated sequences, etc. A therapeutic product may include sterile saline or another pharmaceutically acceptable emulsion and suspension base.

In addition, the kits may include instructional materials containing instructions (e.g., protocols) for the practice of the methods of this invention. While the instructional materials typically comprise written or printed materials they are not limited to such. A medium capable of storing such instructions and communicating them to an end user is contemplated by this invention. Such media include, but are not limited to electronic storage media (e.g., magnetic discs, tapes, cartridges, chips), optical media (e.g., CD ROM), and the like. Such media may include addresses to internet sites that provide such instructional materials.

The present invention also provides for kits for screening for modulators of lung cancer-associated sequences. Such kits can be prepared from readily available materials and reagents. For example, such kits can comprise one or more of the following materials: a lung cancer-associated polypeptide or polynucleotide, reaction tubes, and instructions for testing lung cancer-associated activity. Optionally, the kit contains biologically active lung cancer protein. A wide variety of kits and components can be prepared according to the present invention, depending upon the intended user of the kit and the particular needs of the user. Diagnosis would typically involve evaluation of a plurality of genes or products. The genes typically will be selected based on correlations with important parameters in disease which may be identified in historical or outcome data.

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Example 1: Gene Chip Analysis

Molecular profiles of various normal and cancerous tissues were determined and analyzed using gene chips. RNA was isolated and gene chip analysis was performed as described (Glynne, et al. (2000) Nature 403:672-676; Zhao, et al. (2000) Genes Dev. 14:981-993).

Tables 1A and 1B were previously filed on April 18, 2001 in USSN 60/284,770 (18501-001500US) and on November 29, 2001 in USSN 60/334,370 (18501-001520US)

5	Table 1A					
•	Pkey	ExAcon	UnigenelD	Unigene Title	70% chron/90% NL	70% SQAD/90% NL
	100134	D13264	Hs.49	macrophage scavenger receptor 1	1.61	0.74 3.28
	100780 100971	HG3731-HT4001 J02874	Hs.83213	***Immunoglobulin Heavy Chain, Vdjrc Reg fatty acid binding protein 4; adipocyte	2.68 1.96	0.14
10	101088	L05568	Hs.553	solute carrier family 6 (neurotransmitte	0.79	0.07
-	101102	L07594	Hs.79059	transforming growth factor; beta recepto	2.55	1
	101168	L15388	Hs.211569	G protein-coupled receptor kinase 5	0.88	0.27 0.26
	101277 101330	L38486 L43821	Hs.118223 Hs.80261	microfibrillar-associated protein 4 enhancer of filamentation 1 (cas-like do	0.89 0.59	0.29
15	101336	L49169	Hs.75678	FBJ murine osteosarcoma viral oncogene h	1.15	0.41
	101345	L76380	Hs.152175	calcitonin receptor-like	0.81	0.31
	101678	M62505	Hs.2161	complement component 5 receptor 1 (C5a I	1.31	0.77
	101764 101771	M80563 M81750	Hs.81256 Hs.153837	S100 calcium-binding protein A4 (calcium myeloid cell nuclear differentiation ant	1,44 0,96	0.82 0.45
20	101842	M93221	Hs.75182	mannose receptor, C type 1	1.27	0.37
	102283	U31384	Hs.83381	guanine nucleotide binding protein 11	1.04	0.3
	102363	U39447	Hs.198241	amine oxidase; copper containing 3 (vasc	0.96	0.26
	102507 102698	U52154 U75272	Hs.193044 Hs.1867	potassium inwardly-rectifying channel; s progastricsin (pepsinogen C)	2.81 0.95	3.45 0.23
25	103025	X54131	Hs.123641	protein tyrosine phosphatase; receptor t	1,62	0.21
	103280	X79981	Hs.76206	cadherin 5; VE-cadherin (vascular epithe	0.9	0.41
	103496	Y09267	Hs.132821	flavin containing monooxygenase 2	1.27	0.49
	103541 103554	Z11697 Z18951	Hs.79197 Hs.74034	CD83 antigan (activated B lymphocytes; I caveolin 1; caveolae protein; 22kD	1.86 1.27	1 0.47
30	104212	AB002298	Hs.173035	KIAA0300 protein	1.17	0.16
- •	104691	AA011176	Hs.37744	ESTs	1.08	0.35
	104825	AA035613	Hs.141883	ESTs	0.75	0.27
	104857 104865	AA043219 AA045136	Hs.19058 Hs.22575	ESTs ESTs	2.6 1.23	3.3 0.49
35	104989	AA102098	Hs.118615	ESTs	0.63	0.32
	105729	AA292694	Hs.3807	ESTs; Weakly similar to PHOSPHOLEMMAN PR	0.86	0.34
	105847	AA398606	Hs.32241	ESTs	1.32	0.4
	105894 106490	AA400979 AA451861	Hs.25691 Hs.115537	calcitonin receptor-like receptor activi ESTs; Weakly similar to dipeptidase prec	0.78 1.2	0.28 0.47
40	106536	AA453997	Hs.23804	ESTs	0.82	0.15
	106605	AA457718	Hs.21103	Homo sapiens mRNA; cDNA DKFZp564B076 (fr	0.99	0.07
	106667	AA481086	Hs.16578	ESTS	1.17	0.4
	106773 106797	AA478109 AA478962	Hs.188833 Hs.169943	ESTs ESTs	1.46 1.18	0.43 0.32
45	106844	AA485055	Hs.158213	sperm associated antigen 6	0.98	0.51
	106870	AA487576	Hs.26530	serum deprivation response (phosphatidy)	1.05	0.14
	106954	AA496980	Hs.204038	ESTS	1.25	0.33 0.4
	107054 107292	AA600150 T30407	Hs.14366 Hs.4789	ESTs ESTs; Weakly similar to oxidative-stress	1.11 1.07	2.58
50	107994	AA036811	Hs.165030	ESTs	0.7	0.21
	107997	AA037388	Hs.82223	Human DNA sequence from clone 141H5 on c	1.02	0.48
	108041	AA041552	Hs.61957	ESTs ·	1.44 1.98	0.51 1
	108087 108382	AA045709 AA074885	Hs.40545 Hs.67726	macrophage receptor with collagenous str	1.52	0.72
55	108435	AA078787	Hs.194101	ESTs .	2.53	1.53
	108480	AA081093	Hs.68055	ESTs	1.56	0.48
	109252	AA194830 F01534	Hs.85944 Hs.26981	ESTs . ESTs	2.69 1.19	3.18 0.65
	109550 109613		Hs.27519	ESTs	1.01	0.29
60	109837		Hs.29792	ESTs	0.81	0.15
	109893	H04768	Hs.30484	ESTs	1.44	0.32
	109984 110099	H09594 H16568	Hs.10299 Hs.23748	ESTs ESTs	0.62 1.01	0.14 0.28
	110837	N30796	Hs.17424	ESTs; Weakly similar to semaphorin F [H.	1.1	0.22
65	111247	N69825	Hs.16762	Homo sapiens mRNA; cDNA DKFZp564B2062 (f	1.26	0.26
	111341		Hs.22483	ESTs	1.57 3.96	0.52 1
	111510 111737	R07856 R25410	Hs.16355 Hs.9218	ESTs ESTs	0.97	0.24
	113195	T57112	110.0210	""yc20g11.s1 Stratagene lung (#937210)	1.22	0.35
70	113238	T62979	Hs.189813	ESTs	2.27	0.45
	113540		Hs.16757	ESTS	1.06 1.16	0.22 0.42
	113552 113606		Hs.16026 Hs.17125	ESTs ESTs	1.48	0.42
	113695		Hs.17948	ESTs	1.54	0.28
75	113946	W84753	Hs.37896	ESTs	1.79	0.72
	114251		Hs.21948	ESTs ESTs; Moderately similar to H1 chloride	1.95 1.42	0.25 0.13
	114359 115230		Hs.153483 Hs.182980	ESTs Moderatery similar to A1 cholder	2.62	0.42
00	115230 115279	AA279760	Hs.63671	ESTs	1.79	0.91
80	115566	AA398083	Hs.43977	ESTs	0.86	0.2
	115985 116166		Hs.173233 Hs.202949	ESTs KIAA1102 protein	0.79 2.29	0.04 0.68
	116279	AA486073	Hs.57362	ESTs Piotein	2.27	0.78
	117023	H88157	Hs.41105	ESTs	1.38	0.16
		•				

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	117209	H99959	Hs.42768	ESTs	1.46	0.48
	118901	N90719	Hs.94445	ESTs	1.51	1 0.48
	118981 119073	N93839 R32894	Hs.39288 Hs.45514	ESTs v-ets avlan erythroblastosis virus E26 o	1.34 1.14	0.46
5	119221	R98105	110.1001.1	""yr30g11.s1 Soares fetal liver spleen	1.32	0.53
	119824	W74536	Hs.184	advanced glycosylation end product-speci	1	0.19
	119861	W80715	11- 50200	ESTs; Moderately similar to !!!! ALU SUB	1.83	0.45 0.55
	120041 120132	W92775 Z38839	Hs.59368 Hs.125019	ESTs ESTs; Highly similar to KIAA0886 protein	1.23 0.91	0.37
10	120467	AA251579	Hs.187628	ESTs	1.87	1.91
	121314	AA402799	Hs.182538	ESTs	1.3	0.31
	121643	AA417078	Hs.193767	ESTs	2.31	0.68
	121690 122633	AA418074 AA454080	Hs.110286 Hs.34853	ESTs inhibitor of DNA binding 4; dominant neg	1.47 1.31	0.51 0.63
15	123978	C20653	Hs.170278	ESTs	1.52	0.32
	124214	H58608	Hs.151323	ESTs	0.93	0.35
	124357	N22401		""yw37g07.s1 Morton Fetal Cochlea Homo	1.29	1
	124438 125167	N40188 W45560	Hs.102550 Hs.102541	ESTs ESTs	1.36 1.46	0.7 0.69
20	125174	W51835	Hs.231082	EST	3.07	3.76
	125422	AA903229	Hs.153717	ESTs	1.34	0.3
	125561	AI417667	Hs.22978	ESTS	1.89 0.94	0.63 0.36
	125831 127002	D60988 R35380	Hs.24979	***HUM145B09B Clontech human fetal brain ESTs	3.02	4.06
25	127307	AA369367	Hs.126712	ESTs; Weakly similar to plL2 hypothetica	1.01	0.69
	127609	AA622559	Hs.150318	ESTs	1.21	0.32
	127959 128468	Al302471 D52193	Hs.124292 Hs.56340	ESTs ESTs	2.5 1.13	1 0.33
	128624	AA479209	Hs.102647	ESTs	1.45	0.58
30	128789	AA486567	Hs.105695	ESTs	1.1	0.34
	128798	AF014958	Hs.105938	chemokine (C-C motif) receptor-like 2	1.16	0.55
	128952 129057	R51076 X62466	Hs.107361 Hs.214742	ESTs; Highly similar to Rap2 interacting CDW52 antigen (CAMPATH-1 antigen)	2.04 1.77	2.4 0.73
	129210	AA401654	Hs.202949	KIAA1102 protein	1.11	0.36
35	129240	W24360	Hs.237868	Interleukin 7 receptor	0.91	0.41
	129402 129565	T63781 X77777	Hs.198726	""yc21g01.s1 Stratagene lung (#937210) vasoactive intestinal peptide receptor 1	1.36 0.67	0.43 0.08
	129593	AA487015	Hs.98314	Homo saplens mRNA; cDNA DKFZp586L0120 (f	1.3	0.42
40	129626	AA447410	Hs.11712	ESTs; Weakly similar to IIII ALU SUBFAMI	1.28	0.46
40	129699	AA458578	Hs.12017	KIAA0439 protein; homolog of yeast ubiqu	1.58 1.13	1 0.53
	129898 129958	N48595 L20591	Hs.13256 Hs.1378	ESTs annexin A3	0.81	0.31
	130273	U59914	Hs.153863	MAD (mothers against decapentaplegic; Dr	0.59	0.22
15	130655	N92934	Hs.17409	cysteine-rich protein 1 (intestinal)	1.44	0.76
45	130657 131061	T94452 N64328	Hs.201591 Hs.22567	ESTs ESTs; Moderately similar to HYPOTHETICAL	0.96 1.51	0.42 0.45
	131066	F09006	Hs.22588	ESTs	0.97	0.37
	131263	R38334	Hs.24950	regulator of G-protein signalling 5	2.34	. 2.82
50	131589 131686	U52100 AA157428	Hs.29191 Hs.30687	epithelial membrane protein 2 Grb2-assoclated binder 2	1.2 0.95	0.62 0.38
50	131751	H18335	Hs.31562	ESTs	1,47	0.52
	132430	T23630	Hs.258675	EST	1.86	2.09
	132476 132836	N67192 F09557	Hs.49476 Hs.57929	Homo sapiens clone TUA8 Cri-du-chat regi slit (Drosophila) homolog 3	1.73 0.91	0.58 0.29
55	133120	X64559	Hs.65424	tetranectin (plasminogen-binding protein	0.82	0.2
	133488	D45370	Hs.74120	adipose specific 2	1.29	0.48
	133565	H57058	Hs.204831	ESTs	2.25 1.65	0.57 0.62
	133651 133835	U97105 AA059489	Hs.173381 Hs.76640	dihydropyrimidinase-like 2 ESTs; Highly similar to RGC-32 [R.norveg	1.16	0.34
60	133978	W73859	Hs.78061	transcription factor 21	0.79	0.27
	133985	L34657	Hs.78146	platelet/endothelial cell adhesion molec	0.99	0.28 0.46
	134299 134300	AA487558 U81984	Hs.8135 Hs.166082	ESTs endothelial PAS domain protein 1.	1.02 0.86	0.42
	134323	AA028976	Hs.8175	Homo saplens mRNA; cDNA DKFZp564M0763 (f	1.19	0.27
65	134343	D50683	Hs.82028	transforming growth factor, beta recepto	1.21	0.67
	134417 134561	D87969 U76421	Hs.82921 Hs.85302	solute carrier family 35 (CMP-stalic act adenosine dearninase; RNA-specific; B1 (h	1.28 2.12	1 0.55
	134624	W67147	Hs.8700	deleted in liver cancer 1	2.35	2.74
70	134696	H88354	Hs.8861	ESTs	1.35	0.33
70	134749 134786	L10955 L06139	Hs.89485 Hs.89640	carbonic anhydrase IV TEK tyrosine kinase; endothelial (venous	0.89 0.48	0.2 0.21
	134869	T35288	Hs.90421	ESTs; Moderately similar to IIII ALU SUB	2.14	2.64
	135346	M21056	Hs.992	phospholipase A2; group (B (pancreas)	0.63	0.13
75	100113	D00591	Hs.84746	Chromosome condensation 1 Homo sapiens mRNA for osteoblast specifi	1 0.5	2.15
, 5	100147 100280	D13666 D42085	Hs.136348 Hs.155314	KIAA0095 gene product	0.5 1.02	2 1.39
	100335	D63391	Hs.6793	platelet-activating factor acetylhydrola	1	5.58
	100360	D78335	Hs.75939	Uridine monophosphate kinase	0.91	2.04
80	100372 100486	D79997 HG1112-HT1112	Hs.184339	KIAA0175 gene product TIGR: ras-like protein TC4	0.75 1.09	2.03 1.93
	100559	HG2197-HT2267		"collagen, type VII, alpha 1"	0.97	3.6
	100576	HG2290-HT2386		"calcitonin/alpha-CGRP, alt. transcript	1	1
	100668 100906	HG2981-HT3938 HG4716-HT5158		"TIGR: CD44 (epican, eli. transcript 12 Guanosine 5-Monophosphate Synthase	0.85 1.18	1.9 2.29
85	100930	HG721-HT4827	•	*TIGR: placental protein 14, endometrial	1	1.45

	W	O 02/086	443			
	100960	J00124	Hs.117729	keratin 14 (epidermolysis bullosa simple	0.84	2.6
	101031	J05070	Hs.151738	"Matrix metalloproteinase 9 (gelatinase	0.77	1.52
	101111	L08424	Hs.1619	Achaete-scute complex (Drosophila) homol	1 0 000	1
_	101124	L10343	Hs.112341	"Protease inhibitor 3, skin-derived (SKA	0.62	2.67 1
5	101175	L18920	Hs.36980	"Melanoma antigen, family A, 2"	1 0.74	4.1
	101204	L24203 M19888	Hs.82237 Hs.1076	Ataxia-telangiectasia group D-associated Small proline-rich protein 1B (comifin)	0.85	2.51
	101431 101448	M21389	Hs.195850	keratin 5 (epidermolysis bullosa simplex	. 0.61	8.83
	101511	M27826	Hs.267319	Endogenous retroviral protease	1.03	1.13
10	101526	M29540	Hs.220529	Carcinoembryonic antigen-related cell ad	1.07	4.61
	101548	M31328	Hs.71642	"Guanine nucleotide binding protein (G p	0.97	1.13
	101625	M57293		"Human parathyroid hormone-related pepti	1	1
	101649	M60047	Hs.1690	Heparin-binding growth factor binding pr	1	2.7
1.	101724	M69225	Hs.620	bullous pemphigoid antigen 1 (230/240kD)	1	8.98
15	101748	M76482	Hs.1925	Desmoglein 3 (pemphigus vulgaris antigen	1	2.78
	101759	M80244	Hs.184601	*Solute carrier family 7 (cationic amino	1.07	2.45 1
	101804	M86699	Hs.169840	TTK protein kinase	1 0.74	1.76
	101806	M86757	Hs.112408	S100 calcium-binding protein A7 (psorias "Homo sapiens connexin 26 (GJB2) mRNA, c	1	7
20	101809 101845	M86849 M93426	Hs.78867	*Protein tyrosine phosphatase, receptor-	i	i
20	101851	M94250	Hs.82045	Midkine (neurite growth-promoting factor	1.13	2.6
	102083	U10323	Hs.75117	"Interleukin enhancer binding factor 2,	1.03	1.61
	102154	U17760	Hs.75517	"Laminin, beta 3 (nicein (125kD), kalini	0.94	3.62
	102193	U20758	Hs.313	secreted phosphoprotein 1 (osteopontin;	0.34	4.59
25	102305	U33286	Hs.90073	chromosome segregation 1 (yeast homolog)	1.45	2.97
	102348	U37519	Hs.87539	Aldehyde dehydrogenase B	0.52	2.25
	102581	U61145	Hs.77256	Enhancer of zeste (Drosophila) homolog 2	0.91	2.46
	102610	U65011	Hs.30743	Preferentially expressed antigen in mela	1	3.88
30	102623	U66083	Hs.37110	"Melanoma antigen, family A, 9 (MAGE-9)"	1	1
30	102669	U71207	Hs.29279 Hs.239	Eyes absent (Drosophila) homolog 2 Forkhead box M1	1.06	2.77
	102696 102829	U74612 U91618	Hs.80962	Neurotensin	1	1
	102888	X04741	Hs.7611B	Ubiquitin carboxyl-terminal esterase L1	1.13	2.59
	102913	X07696	Hs.80342	keratin 15	0.7	4.72
35	102915	X07820	Hs.2258	Matrix Metalloproteinase 10 (Stromolysin	1.15	3.35
	102963	X15943	Hs.37058	"Calcitonin/calcitonin-related polypepti	1	1
	103021	X53587	Hs.85266	"Integrin, beta 4"	1.38	2.34
	103036	X54925	Hs.83169	Matrix metalloprotease 1 (interstitial c	1	14.93
40	103058	X57348	Hs.184510	Stratifin	1.25	4.17 1.72
40	103060	X57766	Hs.155324	matrix metalloproteinase 11 (stromelysin	1 1.16	7.38
	103119 103206	X63629 X72755	Hs,2877 Hs,77367	"Cadherin 3, P-cadherin (placental)" monokine induced by gamma interferon	0.71	1.48
	103242	X76342	Hs.389	"Alcohol dehydrogenase 7 (class IV), mu	1	1
	103242	X82693	Hs.3185	"Lymphocyte antigen 6 complex, locus D;	0.92	1.28
45	103478	Y07755	Hs.38991	S100 calcium-binding protein A2	1.05	5.81
	103558	Z19574	Hs.2785	keratin 17	0.65	6.68
	103576	Z26317	Hs.2631	Desmoglein 2	0.79	1.73
	103587	Z29083 ,	Hs.82128	5T4 Oncofetal antigen	1	3.93
50	103594	Z31560	Hs.816	"SRY (sex determining region Y)-box 2, p	0.71	7.23
5 0	103768	AA089997	11- 0407	"ESTs, Highly similar to integral membra	0.99 0.96	1.8 1.29
	104158	AA454908	Hs.8127	KIAA0144 gene product Human DNA sequence from clone 967N21 on	1.23	7.23
	104558 104689	R56678 AA010665	Hs.88959	ESTs	0.96	2.11
	104733	AA019498	Hs.23071	ESTs	1.18	1.88
55	104906	AA055809	Hs.26802	Protein kinese domains containing protei	1.11	3,15
	104978	AA088458	Hs.19322	ESTs; Weakly similar to !!!! ALU SUBFAMI	1.64	2.89
	105012	AA116036	Hs.9329	"Homo saplens mRNA for fis353, complete	1.19	3.91
	105175	AA186804	Hs.25740	ESTs; Weakly similar to unknown (S.cerev	0.9	4.63
CO	105263	AA227926	Hs.6682	ESTs	0.95	2.87
60	105298	AA233459	Hs.26369	ESTs	1 22	1.13 3.01
	105312	AA233854	Hs.23348	S-phase kinase-associated protein 2 (p45	1.32 1.28	2.31
	105719	AA291644	Hs.36793	Hypothetical protein FLJ23188 ESTs	1.20	1
	105743	AA293300 AA411621	Hs.9598 Hs.8895	ESTs; same as BFH6?	0.94	2.04
65	106231	AA429571	Hs.38002	KIAA1355 protein	1.04	1.5
00	106540	AA454607	Hs.38114	Hypothetical protein FLJ11100	1.26	2.26
	106575	AA456039	Hs.105421	ESTs	1	2
	106632	AA459897	Hs.11950	GPI-anchored metastasis-associated prote	0.87	1.32
	106727	AA465342	Hs.34045	Hypothetical protein FLJ20764	0.87	. 1.59
70	106908	AA490237	Hs.222024	Transcription factor BMAL2 (cycle-like f	0.61	1.6
	107059	AA608545	Hs.23044	RAD51 (S. cerevisiae) homolog (E coli Re	0.48	2.67 1.44
	107104	AA609786	Hs.15243	Nucleolar protein 1 (120kD)	1.01 0.97	2.89
	107151	AA621169	Hs.8687 Hs.291904	ESTs; procollagen I-N proteinase Accessory proteins BAP31/BAP29	1.15	3.65
75	107284 107901	S74039 AA026418	Hs.91539	ESTs	0.72	3.44
. 5	107922	AA028028	Hs.61460	lg superfamily receptor LNIR precursor	1	2.48
	107932	AA029317	Hs.18878	Hypothetical protein FLJ21620	1	1
	108695	AA121315	Hs.70823	KIAA1077 protein	0.91	3.53
00	108857	AA133250	Hs.62180	ESTs	1	1
80	108860	AA133334	Hs.129911	ESTs	0.73	7.3
	108990	AA152296	Hs.72045	ESTs	1	1
	109166	AA179845	Hs.73625	*RAB6 Interacting, kinesin-like (rabkine	1	4.55 1.28
	109424	AA227919	Hs.85962	Hyaluronan synthase 3 Whyaluronan synthase 3 Whyaluronan synthase 3	1 1.42	1.28 2
85	109665	F05012	Hs.27027	Hypothetical protein DKFZp762H1311	1.42	2.16
GJ	109970	H09281	Hs.13234	ESTs	1.10	۵. ۱۷

	W	O 02/086	1443			
	110015	H10998	Hs.7164	A disintegrin and metalloproteinase doma	0.84	1.95
	110156	H18957	Hs.4213	ESTs	0.94	1.41
	110561	H59617	Hs.5199	HSPC150 protein similar to ubiquitin-con ESTs; Weakly similar to neogenin [H.sapi	0.91 0.91	3.18 3.13
5	111223 111345	N68921 N89820	Hs.34806 Hs.14559	Hypothetical protein FLJ 10540	1	1.25
	111876	R38239	Hs.293246	*ESTs, Weakly similar to putative p150 [0.83	1.27
,	111902	R39191	Hs.109445	KIAA1020 protein	0.91	0.91
	112244	R51309	Hs.70823	KIAA1077 protein	0.77 1	3.01 1
10	112973 112989	T17271 T23482	Hs.89981	"cDNA FLJ13308 fis, clone OVARC1001436, "Diacylglycerol kinase, zeta (104kD)"	0.55	1.03
10	113047	T25867	Hs.7549	ESTs	0.87	2
	113095	T40920	Hs.126733	ESTs	1	1,,,
	113531	T90345	Hs.16740	Hypothetical protein FLJ11036	0.42 1.17	1.44 1.73
15	113970 114346	W86748 Z41450	Hs.8109 Hs.130489	ESTs "ATPase, aminophospholipid transporter-i	0.86	0.82
13	114407	AA010188	Hs.103305	ESTs	0.8	1.88
	114471	AA028074	Hs.104613	RP42 homolog	1.06	1.34
	114509	AA043551	Hs.101799	KIAA1350 protein	1.82 0.79	2.32 1.49
20	115060 115091	AA253214 AA255900	Hs.198249 Hs.184523	"Gap junction protein, beta 5 (connexin KIAA0965 protein	0.72	1.92
20	115123	AA256642	Hs.236894	*ESTs, High sim to LRP1_hu low density I	0.59	1.97
	115291	AA279943	Hs.122579	ESTS	1	1.25
	115506	AA292537	Hs.45207	Hypothetical protein KIAA1335	1.15 0.5	1.48 3.29
25	115522 115536	AA331393 AA347193	Hs.47378 Hs.62180	ESTs ESTs	1	1
25	115697	AA411502	Hs.63325	Horno sapiens type II membrane serine pro	i	6.53
	115909	AA436668	Hs.59761	ESTs	1	6.98
	115978	AA447522	Hs.69517	Differentially expressed in Fanconi anem	1	2.31 1.68
30	116028	AA452112 AA456968	Hs.42644 Hs.92030	thioredoxin-like ESTs	0.99 1.14	1.8
50	116107 116134	AA460246	Hs.50441	CGI-04 protein	1.11	1.86
	116157	AA461063	Hs.44298	Hypothetical protein	0.99	1.9
	116158	AA461187	Hs.61762	Hypoxia-Inducible protein 2	0.44 0.62	0.86 3.89
35	116335 116483	AA495830 C14092	Hs.87013 Hs.76118	"Homo sapiens cDNA FLJ10238 fis, clone H Ubiquitin carboxyl-terminal esterase L1	1.04	2.36
55	117320	N23239	Hs.211092	LUNX protein; PLUNC(palate lung & nasal	0.51	0.64
	117557	N33920	Hs.44532	Diubiquitin	1.11	2.63
	117693	N40939	Hs.112110	PTD007 protein	0.98 1	1.79 1.43
40	117881 118368	N50073 N64339	Hs.260622 Hs.48956	Butyrate-induced transcript 1 ESTs	0.67	2.86
40	118566	N68558	Hs.42824	Hypothetical protein FLJ10718	1.21	0.83
	118695	N71781	Hs.50081	KIAA1199 see CVA7.doc	0.88	1.63
	119780	W72967	Hs.191381	ESTs; Weakly similar to hypothetical pro	1	1
45	119845 120102	W79920 W95428	Hs.58561 Hs.132927	G protein-coupled receptor 87 *ESTs, Moderately similar to p53 regulat	í	i
73	120102	W95477	Hs.180479	ESTs	0.69	3.07
	120486	AA253400	Hs.137569	Tumor protein 63 kDa with strong homolog	1.08	12.05
	120859	AA350158	Hs.1619	Achaete-scute complex (Drosophila) homol	1	1
50	120880 120948	AA360240 AA397822	Hs.97019 Hs.104650	EST Hypothetical protein FLJ10292	1.04	2.15
	120983	AA398209	Hs.97587	EST	1	1
	121362	AA405500	Hs.97932	Chondromodulin I precursor	1	1
	121369	AA405657	Hs.128791 Hs.293317	CGI-09 protein "ESTs, Weakly similar to JM27 [H.saplens	1 1	1.8 1
55	121791 123005	AA423978 AA479726	Hs.105577	ESTs	i	i
	123044	AA481549	Hs.130881	B-cell CLL/lymphoma 11A (zinc finger pro	0.95	1.88
	123160	AA488687	Hs.284235	ESTs	1.59 1.19	4.98 1.64
	123479 123571	AA599469 AA608956	Hs.135056 Hs.112619	clone RP5-850E9 on chromosome 20 "ESTs, Weakly similar to PQ0109 Purkinje	1.03	1.14
60	123829	AA620697	Hs.112208	XAGE-1 protein	1.39	2.2
	124006	D60302	Hs.108977	ESTs	1	4.85
	124059	F13673	Hs.99769	ESTs Selzure related gene 6 (mouse)-like	1.49 0.76	- 8.62 0.77
	124960 125218	T15386 W73561	Hs.194766 Hs.110024	NADH:ubiquinone oxidoreductase MLRQ subu	1.33	1.77
65	125453	R06041	Hs.18048	"Melanoma antigen, family A, 10"	0.8	1.42
	125759	AA425587	Hs.82226	Glycoprotein (transmembrane) nmb	1.52	2.26
	125972	AA434562	Hs.35406 Hs.270799	"ESTs, Highly similar to unnamed protein EST	1.05 1	2.48 1.95
	125994 126395	H55782 N7Q192	Hs.278956	Hypothetical protein FLJ12929 .	i	1.35
70	126645	Al167942	Hs.61635	STEAP1 (Homo sapiens BAC clone RG041D11	1	2.23
	127221	Al354332	Hs.72365	ESTs	0.73	3.27 1.94
	127479 128192	AA513722 Al204246	Hs.179729	collagen; type X; alpha 1 (Schmid metaph KIAA1085 protein	0.51 1.8	3.16
	128610	L38608	Hs.10247	activated leucocyte cell adhesion molecu	0.89	0.97
75	128777	U46006	Hs.10526	Cysteine and glycine-rich protein 2	1	1
	128924	AA234962	Hs.26557	Plakophilin 3 Solute earrier family 2 (facilitated a)	1.3 0.84	2.97 2.04
	129041 129099	H58873 H50398	Hs.169902 Hs.108660	"Solute carrier family 2 (facilitated g) "ATP-binding cassette, sub-family C (CFT	0.87	1.04
	129404	AA172056	Hs.111128	ESTs	1	1
80	129466	L42583		"Genbank Homo sapiens keratin 6 isoform	0.72	12.67
	129605	S72493 U26727	Hs.115947 Hs.1174	Keratin 16 (focal non-epidermolytic palm "Cyclin-dependent kinase Inhibitor 2A (m	0.92 0.85	1.5 1.93
	129628 130023	X13461	Hs.239600	Calmodulin-like 3	0.84	1.22
~~	130080	X14850	Hs.147097	"H2A histone family, member X"	0.98	1.96
85	130385	AA126474	Hs.155223	stanniocalcin 2	1	1

	\mathbf{w}	O 02/0864	443			
	130410	V01514	Hs.155421	Alpha-fetoprotein	0.63	0.63
	130441	U35835	Hs.301387	"Human DNA-PK mRNA, partial cds"	1.15	3.65
	130482	L32866	Hs.1578	Baculoviral IAP repeat-containing 5 (sur	1 0.92	1.88 1.96
5	130553 130577	AA430032 M35410	Hs.252587 Hs.162	Pituitary tumor-transforming 1 Insulin-like growth factor binding prote	1,17	4.7
-	130627	L23B0B	Hs.1695	Matrix metalloproteinase 12 (macrophage	0.69	4.05
	130800	AA223386	Hs.19574	ESTs; Weakly similar to katanin p80 subu	1.13	2.41
	130939	AA598689	Hs.21400	ESTS	0.8	0.89 1.15
10	131046 131244	X02530 D38076	Hs.2248 Hs.24763	INTERFERON-GAMMA INDUCED PROTEIN PRECURS RAN binding protein 1	1.13	1.85
	131877	J04088	Hs.156346	Topoisomerase (DNA) II alpha (170kD)	1	1
	131927	AA461549	Hs.34780	"Doublecortex: lissencephaly, X-linked (0.81	0.62
	131965	W90145	Hs.35962	ESTs	0.74	3.27
15	131978	D80008	Hs.36232 Hs.211913	KIAA0186 gene product Small proline-rich protein 1A	1 0.69	1 1.43
15	132354 132543	L05187 AA417152	Hs.5101	ESTs; Highly similar to protein regulati	0.79	4.27
	132632	N59764	Hs.5398	guanine-monophosphate synthetase	1	1.08
	132653	U31201	Hs.54451	"laminin gamma2 chain gene (LAMC2), exon	1	1
20	132659	Z75190	Hs.54481	"Low density lipoprotein receptor-relate	0.89	0.89 4.41
20	132710 132758	W93726 W52432	Hs.55279 Hs.56105	"Serine (or cysteine) proteinase inhibit "ESTs, Weakly similar to WDNM RAT WDNM1	0.64 (1.55	2.08
	132767	L05188	Hs.231622	Small proline-rich protein 2B	0.83	1.66
	132816	M74542	Hs.575	Aldehyde dehydrogenase 3	0.55	0.55
25	132990	AA458761	Hs.18387	transcription factor AP-2 alpha (activat	1	3.53 2
23	133070 133282	U69611 U52960	Hs.64311 Hs.286145	"A disintegrin and metalloproteinase dom "SRB7 (suppressor of RNA polymerase B, y	1.16 1	2.7
	133317	AA215299	Hs.70830	U6 snRNA-associated Sm-like protein LSm7	0.95	1.42
	133370	AA156897	Hs.72157	Homo saplens mRNA; cDNA DKFZp564I1922	1.12	2.55
20	133391	X57579	Hs.727	H.sapiens activin beta-A subunit (exon 2	1.65	1.76
30	133832 134032	H03387 Z81326	Hs.241305 Hs.78589	estrogen-responsive B box protein (EBBP) "Serine (or cysteine) proteinase inhibit	1.02 1	1.39 1
	134168	AA398908	Hs.181634		0.95	1.53
	134218	AA227480	Hs.80205	Pim-2 oncogene	1.36	2.48
25	134405	R67275	Hs.82772	"collagen, type XI, alpha 1""	0.76	2.86
35	134453	X70683	Hs.83484	SRY (sex determining region Y)-box 4	1.89 1.82	3.78 4.11
	134470 134645	X54942 U87459	Hs.83758 Hs.167379	CDC28 protein kinase 2 *Cancer/testis antigen (NY-ESO-1, CTAG1,	0.82	0.83
	134781	M17183	Hs.89626	Parathyroid hormone-like hormone	1	1
40	135002	U19147	Hs.272484	Gantigen 6	1	1
40	100040	M97935	11- 0050	AFFX control: STAT1	0.92	1.25 8.5
	101201 101664	L22524 M60752	Hs.2256 Hs.121017	matrix metalloproteinase 7 (matrilysin; H2A histone family; member A	2.92 1	1
	102025	U03911	Hs.78934	mutS (E. coli) homolog 2 (colon cancer;	0.8	1.61
4.5	102031	U04898	Hs.2156	RAR-related orphan receptor A	1	1
45	102221	U24576	11- 75000	LIM domain only 4	1	1 42
	102270 102339	U30255 U37022	Hs.75888 Hs.95577		1.08 0.88	1.43 1.32
	102391	U41668	Hs.77494		1.07	1.58
5 0	103000	X51956	Hs.146580	enolase 2; (gamma; neuronal)	0.91	1.49
50	103395	X94754	Hs.119503		0.89	1.32 1.25
	105638 105726	AA281599 AA292328	Hs.20418 Hs.9754		0.91 0.94	1.48
	114841	AA234722	Hs.55408		0.78	1.56
	115206	AA262491	Hs.186572	ESTs .	1	1
55	115906	AA436616	Hs.82302		0.74 1.1	2.52 1.51
	119132 124163	R49046 H30539	Hs.107911 Hs.189838	ATP-binding cassette; sub-family 8 (MDR/ ESTs	1	1.31
	126487	AA482505	Hs.184601	solute carrier family 7 (cationic amino	1.01	1.46
60	127141	AA307960	Hs.75478		0.85	1.4
60	128034	AA905754	Hs.75103	tyrosine 3-monooxygenase/tryptophan 5-mo	1	1.18 1.5
	128609 128895	AA234365 R37753	Hs.102456 Hs.106985	ESTs	1.7	2
	130199	Z48579	Hs.172028	a disintegrin and metalloprotease domain	1	1
<i>C</i>	130524	U89995	Hs.159234		1	1
65	133000	U24152	Hs.62402 Hs.75426	p21/Cdc42/Rac1-activated kinase 1 (yeast secretogranin II (chromogranin C)	1 1	1
	133658 135047	M25756 AA460466	Hs.93597	ESTs	i	i
	100053	M27830	7.0.0000		0.88	1.53
70	100114	D00596	Hs.82962		0.68	1.86
70	100128	D11094	Hs.61153 Hs.81892		1.29 0.71	2.03 4.26
	100154 100161	D14657 D14694	Hs.77329		1.02	1.56
	100168	D14874	Hs.394	adrenomedullin	0.46	1.17
75	100187	D17793	Hs.78183		1	1.
75	100188	D21063	Hs.57101		0.97	1.4 1.9
	100217 100220	D26600 D28364	Hs.89545		1.13 1.11	1.53
	100287	D43950	Hs.1600	chaperonin containing TCP1; subunit 5 (e	1.13	2.09
90	100297	D49489	Hs.182429	protein disulfide isomerase-related prot	0.92	1.78
80	100330	D55716	Hs.77152		1.07	1.61 1.87
	100355 100364	D78129 D78586	Hs.154868		0.96 1.49	2.46
	100368	D79987	Hs.153479	extra spindle poles; S. cerevisiae; homo	0.59	1.32
05	100398	D84557	Hs.155462		1.08	1.9
85	100438	D87448	Hs.91417	topoisomerase (DNA) II binding protein	1	2.15

	W	O 02/08644	13			
	100455	D87953	Hs.75789	N-myc downstream regulated	0.91	1.48
	100491	HG1153-HT1153	}	Nucleoside Diphosphate Kinase Nm23-H2s	0.99	1.41
	100518	HG174-HT174	_	Desmoplakin I	1.28	3.17
_	100528	HG1828-HT1857		""Nexin, Glia-Derived"	0.68	1.9 5.44
5	100661	HG2874-HT3018		Ribosomal Protein L39 Homolog	1.1 0.8	1.97
	100667	HG2981-HT3127		***Epican, Alt. Splice 11*** Rad2	1.01	2.12
	100830 101061	HG4074-HT4344 K03515	Hs.944	glucose phosphate isomerase	0.91	1.79
	101131	L10838	Hs.167460	splicing factor; arginine/serine-rich 3	1.23	1.87
10	101162	L14595	Hs.174203	solute carrier family 1 (glutamate/neutr	1.35	2.73
	101181	L19686	Hs.73798	macrophage migration inhibitory factor (1.03	1.78
	101183	L19779	Hs.795	H2A histone family; member O	0.57	1.3
	101216	L25876	Hs.84113	cyclin-dependent kinase inhibitor 3 (CDK	0.7	2.2
	101228	L27706	Hs.82916	chaperonin containing TCP1; subunit 6A (0.99	1.99
15	101233	L29008	Hs.878	sorbitol dehydrogenase	0.82	2.11
	101247	L33801	Hs.78802	glycogen synthase kinase 3 beta	1.2	1.91
	101332	L47276		""Homo sapiens (cell line HL-6) alpha t	0.69	2.78
	101342	L76191	Hs.182018	interleukin-1 receptor-associated kinase	1.04	1.84
20	101396	M15796	Hs.78996	proliferating cell nuclear antigen	0.95	3.55 1.5
20	101423	M18391	Hs.89839	EphA1	1 1.21	1.96
	101445	M21259	Hs.1066	small nuclear ribonucleoprotein polypept	0.93	1.6
	101505	M27398	Hs.75692 Hs.12163	asparagine synthetase eukaryotic translation initiation factor	1.19	1.93
	101525 101535	M29536 M30448	Hs.251669	casein kinase 2; beta polypeptide	0.96	1.42
25	101607	M38690	Hs.1244	CD9 antigen (p24)	1.11	1.25
	101624	M55998	110.12.11	*"*Human alpha-1 collagen type I gene, 3	1.17	1.98
	101758	M77836	Hs.79217	pyrroline-5-carboxylate reductase 1	· 1.77	3.45
	101839	M93036	Hs.692	membrane component; chromosomal 4; surfa	0.71	1.45
	101853	M94362	Hs.76084	lamin B2	0.84	1.19
30	101977	S83364		""putative Rab5-interacting protein (cl	0.89	1.9
	101992	U01038	Hs.77597	polo (Drosophia)-like kinase	0.66	1.46
	102009	U02680	Hs.82643	protein tyrosine kinase 9	1.23	3.35
	102012	U03057	Hs.118400	singed (Drosophila)-like (sea urchin fas	0.85	1.88 2.32
25	102039	U05861	Hs.201967	aldo-keto reductase family 1; member C1	0.93 1	4.28
35	102123	U14518	Hs.1594	centromere protein A (17kD) smail nuclear ribonucleoprotein D3 polyp	0.89	1.42
	102130	U15009	Hs.1575 Hs.75823	ALL1-fused gene from chromosome 1q	0.8	2.95
	102148 102210	U16954 U23028	Hs.2437	eukaryotic translation initiation factor	1.01	1.34
	102210	U24389	Hs.65436	lysyl oxidase-like 1	1.15	2.34
40	102260	U28386	Hs.159557	karyopherin alpha 2 (RAG cohort 1; Impor	1.14	2.69
	102330	U35451	Hs.77254	chromobox homolog 1 (Drosophila HP1 beta	1.05	1.7
	102423	U44754	Hs.179312	small nuclear RNA activating complex; po	1.14	2.99
	102455	U48705	Hs.75562	discoidin domain receptor family; member	1.05	2.01
	102499	U51478	Hs.76941	ATPase; Na+/K+ transporting; beta 3 poly	1.27	1.92
45	102522	U53347	Hs.183556	solute carrier family 1 (neutral amino a	0.84	1.31
	102590	U62136		""Homo sapiens enterocyte differentiali	1.11	1.6
	102676	U72514	Hs.12045	putative protein	1.04	2.17 2.28
	102687	U73379	Hs.93002	ubiquitin carrier protein E2-C	0.86 1,12	1.63
50	102704	U76638	Hs.54089	BRCA1 associated RING domain 1 """Human HiV-1 Nef interacting protein (0.9	1.39
50	102781	U83843 U85658	Hs.61796	transcription factor AP-2 gamma (activat	0.98	2.16
	102784 102827	U91327	Hs.6456	chaperonin containing TCP1; subunit 2 (b	0.96	1.62
	102935	X13482	Hs.80506	small nuclear ribonucleoprotein polypept	1.21	4.2
	102972	X16662	Hs.87268	annexin A8	1.25	2.32
55	102983	X17620	Hs.118638	non-metastatic cells 1; protein (NM23A)	1.03	1.83
	103023	X53793	Hs.117950	multifunctional polypeptide similar to S	1.58	5.44
	103038	X54941	Hs.77550	CDC28 protein kinase 1	1.32	3.79
	103075	X59543	Hs.2934	ribonucleotide reductase M1 polypeptide	1.11	2.58
C O	103168	X68314	Hs.2704	glutathione peroxidase 2 (gastrointestin	0.75	3.05
60	103185	X69910	Hs.74368	transmembrane protein (63kD); endoplasmi	1.01 0.95	1.97 1.72
	103212	X73874	Hs.2393	phosphorylase kinase; alpha 1 (muscle)	0.97	1.77
	103223	X74801	Hs.1708 Hs.3155	chaperonin containing TCP1; subunit 3 (g casein; alpha	1	1
	103260 103262	X78416 X78565	Hs.204133	hexabrachion (tenascin C; cytotactin)	1.23	3.09
65	103202	X85373	Hs.77496	smail nuclear ribonucleoprotein polypept	1.12	2.25
05	103364	X90872	Hs.75854	SULT1C sulfotransferase	2.85	4.62
	103375	X91868	Hs.54416	sine oculis homeobox (Drosophila) homolo	1	2.48
	103391	X94453	Hs.114366	pyrroline-5-carboxylate synthetase (glut	1	1.53
	103404	X95586	Hs.78596	proteasome (prosome; macropain) subunit;	0.92	1.53
70	103437	X98260	Hs.82254	M-phase phosphoprotein 11	0.92	1.54
	103448	X99133	Hs.204238	lipocalin 2 (oncogene 24p3)	0.55	0.96
	103605	Z35402	Hs.194657	cadherin 1; E-cadherin (epithelial)	1.32	2.51
	103646	Z68228	Hs.2340	junction plakoglobin	0.88	1.28
75	103658	Z74615	Hs.172928	collagen; type I; alpha 1	1.06	2.98 4.66
75	103774	AA092898	Hs.92918	ESTs; Weakly similar to R07G3.8 [C.elega	1.88 0.87	2.17
	104261	AF008442	Hs.5409	RNA polymerase I subunit ESTs; Weakly similar to R27090_2 [H.sapi	1.4	2.17
	104276	C02193	Hs.85222	KIAA0956 protein	1.15	1.68
	104289 104434	C16281 L02870	Hs.75478 Hs.1640	collagen; type VII; alpha 1 (epidermolys	1.04	1.49
80	104453	M19169	Hs.123114	cystetin SN	0.38	0.76
	104611	R98280	Hs.125845	ribulose-5-phosphate-3-epimerase	1.08	2.25
	104758	AA024661	Hs.7010	ESTs; Weakly similar to ACYL-COA DEHYDRO	1.14	1.65
	105114	AA156532	Hs.11801	adenosine A2b receptor pseudogene	0.91	1.38
0.5	105132	AA159501	Hs.247280	HBV associated factor	1.08	1.7
85	105174	AA186613	Hs.34744	ESTs	0.95	2.05

	W	O 02/086	5443			
	105280	AA232215	Hs.14600	ESTs	1	1.4
	105344	AA235303	Hs.8645	ESTs	0.72	2.02
	105516	AA257971	Hs.21214	ESTs	1.35	3.56 1.82
5	105621	AA280865	Hs.6375 Hs.15202	Homo sapiens mRNA; cDNA DKFZp564K0222 (f	1.23 0.98	1.28
,	105698 105705	AA287393 AA290767	Hs.101282	ESTs; Weakly similar to oligodendrocyte- Homo saplens mRNA; cDNA DKFZp434B102 (fr	0.92	1.32
	105703	AA292098	Hs.22934	ESTs; Weakly similar to ZINC FINGER PROT	0.99	1.41
	105782	AA350215	Hs.21580	ESTs	1	1
	105799	AA372018	Hs.24743	ESTs	1.08	1.78
10	105807	AA393803	Hs.16869	ESTs; Moderately similar to COLLAGEN ALP	0.95	1.34
	105891	AA400768	Hs.26662	ESTs; Weakly similar to tumor necrosis f	0.87	2.25
	105936	AA404338	11- 00000	ESTs	1.14 1	1.46 1
	106069 106103	AA417741	Hs.29899 Hs.12094	ESTs; Weakly similar to ZINC FINGER PROT ESTs	1.04	1.44
15	106140	AA421104 AA424524	Hs.14912	KIAA0286 protein	1.23	2.11
15	106149	AA424881	Hs.256301	ESTs	0.83	1.48
	106154	AA425304	Hs.6994	ESTs	0.77	2.05
	106182	AA426609	Hs.10862	ESTs	0.74	2,23
20	106220	AA428582	Hs.32196	ESTs; Moderately similar to metargidin p	0.97	1.99
20	106228	AA429290	Hs.17719	ESTs	0.99 0.95	1.54 2.09
	106318	AA436570	Hs.9605	pre-mRNA cleavage factor Im (25kD)	0.98	2.66
	106341 106432	AA441798 AA448850	Hs.5243 Hs.17138	ESTs; Moderately similar to ptL2 hypothe ESTs	0.95	1.93
	106474	AA450212	Hs.42484	Homo sapiens mRNA; cDNA DKFZp564C053 (fr	1	1
25	106483	AA451676	Hs.30299	IGF-II mRNA-binding protein 2	1.4	2.29
	106599	AA457235	Hs.12842	ESTs; Moderately similar to non-function	1	1.82
	106611	AA458904	Hs.26267	ESTs; Weakly similar to torsInA [H.saple	1.49	2.78
	106654	AA460449	Hs.3784	ESTs; Highly similar to phosphoserine am	1	1.4
30	107076	AA609145	Hs.21143	ESTs; Weakly similar to fos39554_1 [H.sa	1.11 1	1.49 1.03
30	107115 107129	AA610108 AA620553	Hs.27693 Hs.4756	ESTs; Highly similar to CGI-124 protein flap structure-specific endonuclease 1	1.13	3.63
	107159	AA621340	Hs.10600	ESTs; Weakly similar to ORF YKR081c [S.c	1.05	2.09
	107444	W28391	Hs.5181	proliferation-associated 2G4; 38kD	1.18	1.9
~ -	107481	W58247	Hs.27437	Homo sapiens kinesin superfamily motor K	0.99	2.74
35	107516	X56597	Hs.99853	fibrillarin	0.94	1.77
	107529	Y12065	Hs.5092	nucleolar protein (KKE/D repeat)	1.05	2.29
	107531	Y13936	Hs.17883 Hs.173100	protein phosphatase 1G (formerly 2C); ma	1.06 1.03	1.62 1.4
	107801 107957	AA019433 AA031948	Hs.57548	ESTs ESTs	0.95	1.46
40	108565	AA085342	Hs.1526	ATPase; Ca++ transporting; cardiac muscl	0.59	1.35
	108780	AA128561	Hs.117938	collagen; type XVII; alpha 1	1	7.63
	108828	AA131584	Hs.71435	DKFZP564O0463 protein	1.33	2.56
	109060	AA160879	Hs.241551	chloride channel; calcium activated; fam	0.67	1.42
15	109112	AA169379	Hs.72865	ESTs	1.03	2.31
45	109344	AA213696	Hs.86559	poly(A)-binding protein-like 1 ESTs; Weakly similar to REGULATOR OF MIT	0.97 0.76	1.55 1.87
•	109412 110780	AA227145 N23174	Hs.209473 Hs.22891	solute carrier family 7 (cationic amino	0.9	0.95
	110958	N50550	Hs.24587	signal transduction protein (SH3 contain	1.17	2.26
	111018	N54067	Hs.3628	mitogen-activated protein kinase kinase	1.21	1.85
50	111337	N79612	Hs.16607	ESTs; Highly similar to Myosin heavy cha	1	1.45
	112305	R54822	Hs.26244	ESTs	1	1
	112401	R61279	Hs.237536	ESTs; Weakly similar to F25B5.3 [C.elega EST	1.24 1.56	1.64 1.96
	112853 112869	T02843 T03313	Hs.4351 Hs.4747	dyskeralosis congenita 1; dyskerin	1.03	1.57
55	112992	T23513	Hs.7147	ESTs	1	1
	113048	T25895	Hs.184008	ESTs; Weakly similar to RNA-binding prot	1.37	2.26
	113063	T32438	Hs.5027	ESTs	1	1
	113179	T55182	Hs.152571	ESTs; Highly similar to IGF-II mRNA-bind	1.33	2.7
60	113573	T91166	Hs.15990	ESTs	0.76 0.79	1.47 1.51
00	113811 114086	W44928 Z38266	Hs.4878 Hs.12770	ESTs Homo saplens PAC clone DJ0777O23 from 7p	0.9	1.34
	114587	AA070827	Hs.180320	ESTs; Weakly similar to GOLGI 4-TRANSMEM	1.02	1.76
	114846	AA234929	Hs.44343	ESTs	1.32	2.36
~~	114964	AA243873	Hs.82184	ring finger protein 3	1.1	1.84
65	115047	AA252627	Hs.22554	homeo box 85	1.01	2.36
	115166	AA258409	Hs.198907	myelin protein zero-like 1 hypothetical protein	1.05 1.52	2.31 2.52
	115167 115239	AA258421 AA278650	Hs.43728 Hs.73291	ESTs; Weakly similar to similar to the b	0.7	2.57
	115278	AA279757	Hs.67466	ESTs; Weakly similar to BACN32G11.d [D.m	1.14	2.12
70	115652	AA405098	Hs.38178	ESTs	0.82	4.67
• -	115875	AA433943	Hs.43946	ESTs; Weakly similar to Weak similarity	1.2	1.98
	116004	AA449122	Hs.76086	ESTs; Highly similar to small zinc finge	0.96	1.31
	116121	AA459254	Hs.48855	ESTs	0.97	1.55
75	116129	AA459956	Hs.49163	ESTs; Highly similar to putative ribonuc	1.08 0.8	2.73 1.57
75	116190	AA464963 AA490494	Hs.67776 Hs.65403	ESTs ESTs	1.37	2.65
	116312 116732	F13779	Hs.65403 Hs.165909	ESTs	0.92	1.8
	117602	N35020	Hs.44685	ESTs; Weakly similar to GOLIATH PROTEIN	1.15	1.84
	117950	N51394	Hs.75478	KIAA0956 protein	1.04	2.36
80	117992	N52000	Hs.172089	Homo sapiens mRNA; cDNA DKFZp586B0222 (f	0.62	1.29
	118785	N75386	Hs.111867	GLI-Kruppel family member GLI2	1	1
	119717	W69134	Hs.57987	ESTs ESTs	1 0.78	1.4 1.77
	119814 120128	W74069 Z38499	Hs.58350 Hs.91448	MKP-1 like protein tyrosine phosphatase	0.86	1.46
85	120126	Z98443	Hs.86366	ESTs	0.83	2.01

12065 A-325804 Hs. 1578 1578		w	∩ 02/086	1113			
121366 AA490266 H. 46.97031 ESTR, Weakly similar to Smillar to phyth 0.98 1.3. 121376 AA401246 H. 46.97031 ESTR, Weakly similar to Smillar to phyth 0.99 1.89 121377 AA411446 H. 46.20885 ESTR, Weakly similar to SOU(UM-AND 0.91 1.89 121780 AA421326 H. 46.20895 ESTR 0.001147 H. 46.101147 H. 46.101147 ESTR 0.001147 H. 46.101147 H. 46.101147 ESTR 0.001147 H. 46.101147 H. 46.101147 H. 46.101147 ESTR 0.001147 H. 46.101147 H. 46.1					anonhosis inhibitor 4 (survivin)	0.74	1.64
121326 AA40426 Hs.97031 ESTR, Weakly similar to Strinker to phylo							
5 121497 AA42208 Ha.12660 ESTs							
12781 AA42208 Hs.12660 EST8 cycleriums PS40 family member predicted 1.07 1.54 A422172 Hs.98470 gpp junction protein; bela 2, 2805 (com 0.34 1.43 1.43 1.43 1.43 1.43 1.43 1.43 1	5						
121814 AA425772 h. h. 98470 cyto-hrome P506 family member produkted 1.07 1.54 1.54 1.20598 AA43377 h. h. 98478 EST 1.30 2.23 1.20598 AA43377 h. h. 98478 EST 1.30 2.23 1.20598 AA43377 h. h. 98478 EST 1.30 2.23 1.20598 AA43377 h. h. 98498 EST 1.30 2.23 1.20598 AA43377 h. h. 98498 EST 1.30 2.23 1.20598 AA43378 h. h. 198591 EST 1.30 2.23 1.20598 AA43378 h. h. 198591 EST 1.30598 A. 600847 h. h. 170913 EST 1.30598 AA621826 h. h. 105514 EST 1.30573 AA661826 h. h. 105514 EST 1.30573 AA681823 h. h. 105514 EST 1.30573 AA681825 H. h. 105514 EST A. 105514 ES	3						
120599 AA443772 He.89978 EST 1 1.30 2.23							1.54
122358		121844					
122514 AA44377	1Λ						
122991 AA452356 Hs. 991911 ESTs Weekly similar to MRJ [N. septene] 2.28 2.93 12398 AA605631 Hs. 199516 ESTs 1 1.93	10						
123398 AAS21265 Hs. 105514 EST6 1 1.93						2.28	
123737 AA609471 1-1, 1-1, 1-1, 1-1, 1-1, 1-1, 1-1, 1							
128673 AA698471 Hs.112712 ESTs	15						
124907 N24006 He.99348 He	13						
12444							
2575 W25958 Hs. 81634 Hs. 82163 1599 156 156 157 159							
125769 A3262847 Hs. 82128 T34 concletal trophoblast glycoprolein 1.55 6.76 6.76	20						
128852 H09290	20						
128037 M85772							
1921/4 N78770 1.82 1.8							
128414 N78770	25						
126737 AA488132 Hs.172612 127873 AA479854 Hs.172612 128264 AA179854 Hs.172612 128264 Hs.2822 Hs.101042 A5691734 Hs.170311 128218 Hb2682 Hs.8322 Hs.101042 128268 X6973 Hs.247568 128268 X69673 Hs.247568 128268 X69673 Hs.247568 128268 X6973 Hs.101850 128269 X7939 Hs.103834 Hs.101850 128731 A328993 Hs.101850 128251 A348527 Hs.101850 128251 A348528 Hs.101850 128251 A32850	20						
128928		126737					
127432							
128218 M205682 Hs. 199189 ESTs Moderately similar to recombination 1.24 2.09 1.2568 1.2568 M31433 Hs. 101958 Has. 24758 Earlyste kinase 3 1.23 3.48 1.23 3.48 1.25828 1.25828 (14037 Hs. 2451978 ESTs 1.22 1.9 1.22714 1.2273 A.322893 Hs. 104585 ESTs 1.1 1.73 1.2273 A.322893 Hs. 104585 ESTs 1.1 1.73 1.2273 A.322893 Hs. 104585 ESTs 1.1 1.73 1.2273 A.322893 Hs. 104585 ESTs 1.2550 Hs. 1.05465 Hs. 108523	30						
128568 X66973 Hs.247568 aderlysis kinase 3 1.23 3.48	50					1.24	
12858							
128282 C.14037							
128691 W27939	35						
129733 AA328993			W27939	Hs.103834			
128781							
129052							
129241	40				ribosomal protein S11		
12965							
45 129720 AA401348 Hs. 17999 ESTE							
129720							
12885	45	129720	AA476582				
130069 AA055896 Hs. 146428 collagen; type V; alpha 1 1.17 1.98 1.30405 H88339 Hs. 155398 Hs. 155398 Hs. 155398 Hs. 165398 Hs. 165398 Hs. 164070 Hs. 174070 130599 M91670 Hs. 174070 Hs. 174070 Hs. 174070 130697 Jo4093 Hs. 2056 UDP glycosyltransferase 1 1 4.8 1.07 1.66 UDP glycosyltransferase 1 1 4.8 1.31083 US6661 Hs. 2277 Hs. 2277 US48ily similar to NADH-CYTOCHROME 0.93 1.05 US48ily similar to NADH-CYTOCHROME 0.93 1.05 US4114 US48ily similar to NADH-CYTOCHROME 0.93 1.05 US48ily similar to Mportal t							
130405							
130559 M91670 Hs.174070 Ubiquilin carrier protein 1.07 1.66 130687 Jo4093 Hs.2056 UDP glycosyltransferase 1 1 4.8 131009 AA063596 Hs.2214 ESTs; Weakly similar to NADH-CYTOCHROME 0.93 1.05		130405	H88359	Hs.155398	nuclear factor (erythroid-derived 2)-lik		
130667 J04093	50						•
131009							
131083 U66661					ESTs; Weakly similar to NADH-CYTOCHROME		
131091 T35341 Hs.22880 ESTs; Highly similar to dipeptidy) pept 1.28 1.98	55						
131144 C14412 Hs.23528 ESTs; Highly similar to HSPC038 protein 1.43 2.08 131148 C00038 Hs.23579 ESTs 0.88 3.38 131164 Y00503 Hs.182265 keralin 19 1.19 2.77 1.19 2.77 1.19 1.19 2.77 1.19 1.19 2.77 1.19 1.19 2.77 1.19 1.19 2.77 1.19 1.19 2.77 1.19 1.19 2.77 1.18 1.19 2.77 1.18 1.19 2.77 1.18 1.19 2.77 1.18 1.18 1.19 2.77 1.18 1.18 1.18 1.18 1.18 1.18 1.18 1.18 1.18 1.18 1.18 1.18 1.1	22						
131164 Y00503	•						2.06
60 131185 M25753 Hs.23960 cyclin B1							
131219 C00476	60						
131454 AA455896	00						2.96
131689		131454	AA455896				
65 131692 D50914 Hs.30738 KIAA0124 protein 1.55 2.39 131786 AA135554 Hs.32125 ESTs 1 1.33 131843 AA195893 Hs.184062 ESTs; Moderately similar to putative Rab 0.83 1.63 131860 U02082 Hs.334 Oncogene TIM 1.08 2.2 131803 AA481723 Hs.3463 ribosomal protein S23 1.23 1.24 131903 AA481723 Hs.3463 deleted in oral cancer (mouse; homolog) 0.91 1.18 131958 AA093998 Hs.35120 replication factor C (activator 1) 4 (37 1 2.8 131958 AA093998 Hs.3566 ESTs; Highly similar to phosphorylation 0.87 1.36 132064 W42508 Hs.3593 ESTs 1 1.25 132040 AA146843 Hs.172894 BH3 interacting domain death agonist 1 1.55 132065 D82226 Hs.211594 proteasome (prosome; macropain) 26S subu 0.89 1.27 132112 <td></td> <td></td> <td></td> <td></td> <td></td> <td></td> <td></td>							
131786	65						
131860 U02082 Hs.334 Oncogene TIM 1.08 2.2				Hs.32125	ESTs		
131884	,						
70 131903 AA481723 Hs.3436 deleted in oral cancer (mouse; homolog) 0.91 1.18 131945 M87339 Hs.35120 replication factor C (activator 1) 4 (37 1 2.8 131958 AA093998 Hs.3566 ESTs; Highly similar to phosphorylation 0.87 1.36 131964 W42508 Hs.3593 ESTs 1 1.25 132001 J00277 Hs.37003 v-He-ras Harvey rat sarcome viral oncoge 1.12 1.43 132065 D82226 Hs.211594 BH3 Interacting domain death agonist 1 1.55 132109 AA599801 Hs.40098 ESTs 1 1.05 132112 AA150661 Hs.40154 jumonjl (mouse) homolog 0.99 1.44 132123 AA447123 Hs.250705 ESTs 1.06 2.46 132180 AA4695569 Hs.418 fibroblast activation protein; alpha; se 1.08 2.45 132309 AA460917 Hs.2780 jun D proto-oncogene 1.16 1.8 132318							
131958 AA093998 Hs.3566 ESTs; Highly similar to phosphorylation 0.87 1.36 131964 W42508 Hs.3593 ESTs 1 1 1.25 132001 J00277 Hs.37003 v-He-ras Harvey rat sarcoma viral oncoge 1.12 1.43 132065 D82226 Hs.211594 proteasome (prosome; macropain) 26S subu 0.69 1.27 132109 AA599801 Hs.40098 ESTs 1 1 1.05 132112 AA150661 Hs.40154 jumonji (mouse) homolog 0.99 1.44 132123 AA447123 Hs.250705 ESTs 1.06 2.46 132162 H89551 Hs.41241 ESTs 1.08 2.46 132162 AA095689 Hs.418 fibroblast activation protein; alpha; se 1.02 4.56 132309 AA460917 Hs.2780 jun D proto-oncogene 1.16 1.8 1.32187 AA253448 Hs.46577 ESTs 0.8 1.26 1.32187 AA253448 Hs.45657 ESTs 0.8 1.26 1.32180 AA253330 Hs.53444 adentor-related protein complex 1; gamma 0.5 1.49	70				deleted in oral cancer (mouse; homolog)	0.91	
131964 W42508 Hs.3583 ESTs 1 1.25							
75 132040 AA148843 Hs.172894 BH3 Interacting domain death agonist 1 1.55 132040 AA148843 Hs.172894 BH3 Interacting domain death agonist 1 1.55 132165 D82226 Hs.211594 proteasome (prosome; macropain) 26S subu 0.89 1.27 132107 AA599801 Hs.40098 ESTs 1 1.05 132112 AA150661 Hs.40154 jumonji (mouse) homolog 0.99 1.44 132123 AA447123 Hs.250705 ESTs 1.06 2.46 132162 H89551 Hs.41241 ESTs 1.08 2.45 132180 AA405569 Hs.418 fibroblast activation protein; alpha; se 1.02 4.56 132309 AA460917 Hs.2780 jun D proto-oncogene 1.16 1.8 132311 AA253330 Hs.5344 Adenot-related protein complex 1; gamma 0.5 1.49							
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132109	75	132040					
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80 132123 AA447123 Hs.260705 ESTs 1.06 2.46 132162 H89551 Hs.41241 ESTs 1.08 2.48 132180 AA405569 Hs.418 fibroblast activation protein; alpha; se 1.02 4.56 132309 AA460917 Hs.2780 Jun D proto-oncogene 1.16 1.8 132371 AA235448 Hs.46677 ESTs 0.8 1.26 13238 AA253330 Hs.5344 adentor-related protein complex 1; gamma 0.5 1.49							
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132309 AA460917 Hs.2780 Jun D proto-oncogene 1.16 1.8 132371 AA235448 Hs.46677 ESTs 0.8 1.26 132618 AA253330 Hs.5344 adentor-related protein complex 1; gamma 0.5 1.49	80	132162	H89551	Hs.41241			
132371 AA235448 Hs.45677 ESTs 0.8 1.26							
132618 AA253330 Hs.5344 adentor-related protein complex 1; gamma 0.5 1.49					ESTs	8.0	1.26
OJ 132738 U68019 Hs.211578 MAD (mothers against decapentaplegic; Lir 1.21 1.81	05	132618	AA253330	Hs.5344	adaptor-related protein complex 1; gamma		
	03	132736	U68019	Hs.211578	MAD (mothers against decapentaplegic; Dr	1.21	1.01

	W	O 02/086	443			
	132771	AA488432	Hs.56407	phosphoserine phosphatase	1	1.3
	132833	U78525	Hs.57783	eukaryotic translation initiation factor	0.91	1.43
	132922	T23641	Hs.6066	KIAA1112 protein	1.16	1.53
	132959	AA028103	Hs.61472	ESTs; Weakly similar to unknown [S.cerev	1.02	1.88
5	132994	AA505133	Hs.7594	solute carrier family 2 (facilitated glu	0.72	2.97
•	133005	C21400	Hs.103329	KIAA0970 protein	0.88	1.34
	133065	X62535	Hs.172690	diacylglycerol kinase; alpha (80kD)	0.93	1.23
	133083	N70633	Hs.6456	chaperonin containing TCP1; subunit 2 (b	1.14	1.76
	133086	L17131	Hs.139800	high-mobility group (nonhistone chromoso	0.97	1.43
10	133134	T89703	Hs.65648	RNA binding motif protein 8	1.1	1.8
	133195	AA350744	Hs.181409	KIAA1007 protein	2.29	2.69
	133313	AA249427	Hs.70704	ESTs	1.07	1.68
	133331	T62039	Hs.158675	ribosomal protein L14	0.85	1.18
	133438	D13370	Hs.73722	APEX nuclease (multifunctional DNA repal	0.91	1.45
15	133445	T99303	Hs.73797	guanine nucleotide binding protein (G pr	0.94	1.68
	133483	X52426	Hs.74070	keratin 13	0.85	1.14
	133492	L40397	Hs.74137	transmembrane trafficking protein	1.1	1.69
	133504	W95070	Hs.74316	desmoplakin (DPI; DPII)	0.7	6.21
	133517	X52947	Hs.74471	gap junction protein; alpha 1; 43kD (con	0.95	1.3
20	133540	D78151	Hs.74619	proteasome (prosome; macropain) 26S subu	0.91	1.25
	133594	L07758	Hs.172589	nuclear phosphoprotein similar to S. cer	0.84	1.29
	133627	U09587	Hs.75280	glycyl-tRNA synthetase	1.09	1.99
	133671	T25747	Hs.75471	zinc finger protein 146	1.02	1.5
	133859	U86782	Hs.178761	26S proteasome-associated pad1 homolog	1.11	3.33
25	133865	F09315	Hs.170290	discs; large (Drosophila) homolog 5	1.84	6.7
	133913	W84712	Hs.7753	calumenin	1.15	1.86
	133963	L34587	Hs.184693	transcription elongation factor B (SIII)	1.3	1.91
	133982		Hs.207251	nucleolar autoantigen (55kD) similar to	1.3	1.99
20	134100	L07540	Hs.171075	replication factor C (activator 1) 5 (36	0.72	1.65
30	134110	U41060	Hs.79136	LIV-1 protein; estrogen regulated	1.04	1.62
	134158	U15174	Hs.79428	BCL2/adenovirus E1B 19kD-interacting pro	1	1.55
	134161	U97188	Hs.79440	IGF-II mRNA-binding protein 3	0.82	1.95
	134193	F09570	Hs.7980	ESTs	0.98	1.48
25	134367	X54199	Hs.82285	phosphoribosylglycinamide formyltransfer	1	2.8
35	134402	U25165	Hs.82712	fragile X mental retardation; autosomal	1.26	2 1.47
	134457	D86963	Hs.174044	dishevelled 3 (homologous to Drosophila	1 0.94	1.57
	134469	X17567	Hs.83753	small nuclear ribonucleoprotein polypept	0.94 1.2	2.64
	134498	M63180	Hs.84131	threonyl-tRNA synthetase eukaryotic translation initiation factor	0.84	1.36
40	134501	W84870	Hs.211568	replication protein A1 (70kD)	1.7	2.93
40	134507	M63488	Hs.84318 Hs.85215	Deleted in split-hand/split-foot 1 regio	1.46	2.73
	134548 134599	U41515 X99226	Hs.86297	Fanconi anemia; complementation group A	1.36	2.22
	134692	R73567	Hs.8850	a disintegrin and metalloproteinase doma	0.77	1.64
	134693	N70361	Hs.8854	ESTs	1.09	1.82
45	134806	Z49099	Hs.89718	spermine synihase	0.98	1.35
7.5	134821	Z34974	Hs.198382	plakophilin 1 (ectodermal dysplasia/skin	0.99	1.4
	134864	Y08999	Hs.90370	actin related protein 2/3 complex; subun	0.95	1.42
	134914	U29615	Hs.91093	chitinase 1 (chitotriosidase)	1.16	1.29
	134953	L10678	Hs.91747	profilin 2	0.95	1.76
50	134993	AA282343	Hs.9242	purine-rich element binding protein B	0.98	1.73
	135051	C15324	Hs.93668	ESTs	1.35	2.11
	135158	U51711		Human desmocollin-2 mRNA; 3' UTR	0.86	1.16

Table 1B shows the accession numbers for those pkeys in Table 1A lacking unigenelD's. For each probaset we have listed the gene cluster number from which the oilgonucleotides were designed. Gene clusters were compiled using sequences derived from Genbank ESTs and mRNAs. These sequences were clustered based on sequence similarity using Clustering and Alignment Tools (DoubleTwist, Oakland California). The Genbank accession numbers for sequences comprising each cluster are listed in the Accession column.

	Pkey: Unique Eos probeset identifier number
60	CAT number: Gene cluster number
	Accession: Genbank accession numbers

	Pkey	CAT	Accessions
65	100661 100667	23182_1 26401_3	BE623001 L05098 AA383604 AW966416 N53295 AA460213 AW571519 AA603655 L05424 X56794 S66400 X55150 W60071 AW351820 X55938 M83326 BE005289 BE070059 M83324 BE005248 BE089717 BE181648 BE069700 AW606203 BE069721 AW382138 AW803776 BE463954 BE005334 BE005274 T27386 AA932714 AA972695 AW377728 AI632506 T29068 AJ783934 AW377727 BE163715 AL047291 AA279047 AA523003 BE008048 BE440141 W23614 BE090519 BE092193 N29181 N20358 N44153
70			BE546944 T69231 AW377441 AA907406 H50799 AW051416 Al420712 BE620922 Al279161 AA992549 W47198 BE005241 Al342698 H50700 Al969974 Al863855 AA374490 AW130675 Al950633 AA146687 H99482 X55150 BE005414 BE005339 N28294 Al673068 Al887890 AW804171 Al675961 AW804172 AA778841 AL048050 Al127757 Al095568 AW204965 AW468978 W31898 Al052595 Al278771 BE464018 Al081503 Al824196 AA513211 AA411062 AW084376 N48752 AA703209 N35580 AW059918 AA054563 Al280942 T27619 BE621435 N66010 AW589527 Al160414 AA283090 AA962536 H82726 W52115 W45432 W60433 AA577548 AA146714 BE150994 AA054615 AW796025 AW382768 BE565671 C00444 AA054555
75	100668	26401_3	L05424 X56794 S66400 X55150 W60071 AW351820 X55938 M83326 BE005289 BE070059 M83324 BE005248 BE069717 BE181648 BE069700 AW606203 BE069721 AW382138 AW803776 BE463954 BE005274 T27386 AA932714 AA972695 AW377727 BA1632506 T29068 AJ763934 AW377727 BE163715 AL047291 AA279047 AA523003 BE008048 BE440141 W23614 BE090519 BE092193 N29181 N20358 N44153 BE546944 T69231 AW377441 AA907406 H50799 AW051416 Al420712 BE620922 Al279161 AA992549 W47198 BE005241 Al342695 H50700
80			A1969974 A1863855 AA374490 AW130675 A1950633 AA146687 H99482 X55150 BE005414 BE005339 N28294 A1673068 A1887890 AW804171 A1675961 AW804172 AA778841 AL048050 A1127757 A1095568 AW204965 AW468978 W31898 A1052595 A1278771 BE464018 A1081503 A1824196 AA513211 AA411062 AW084376 N48752 AA703209 N35580 AW059918 AA054563 A1280942 T27619 BE621435 N66010 AW589527 A1160414 AA283090 AA962536 H82726 W52115 W45432 W60433 AA577548 AA146714 BE150994 AA054615 AW796025 AW382768 BE565571 C00444 AA054555
85	101332	25130_1	J04088 RM_001067 AF071747 AJ011741 N85424 AL042407 AA218572 BE296748 BE083981 AL040877 AW499918 AW675045 H17813 BE081283 AA670403 AW504327 BE094229 AA104024 AI471482 AI970337 AA737616 AI827444 AW003286 AI742333 AI344044 AI765634

	***	0 02/000	Al948838 AW235336 AW172827 AA095289 BE046383 Al734240 W18699 Al660329 Al289433 AA933778 AW469242 AA468838 AA806983
5			AJ848838 AW235336 KM77827 AAU35289 BEU46363 AI734240 W16699 A1660529 A169533 A16374 AW402322 AW6504319 A1221834 A1337434 AA625873 W78031 BE206307 AA550803 A1743147 A1990075 AA948274 AA129533 A1635399 AA605313 A1624669 AW594319 A1221834 A1337434 AA307706 BE550282 A1760467 A1630836 A1221821 AW674314 AW078889 A1953732 A1686969 A1186928 AW074595 A1127486 A1079644 A1910815 H17814 AA310903 AW137854 T19279 AA026682 AA306035 AW383390 AW383389 AW383422 AW383427 AW383395 H09977 AA306247 AA352501 AW403639 F05421 AA224473 AA305321 H93904 AA089612 AW391543 AW402915 AW173382 AW402701 AW403113
3			A3U6247 AA325011 AW40U639 FU3421 AA224473 AA305321 FISSIGN ANA069112 AW379154 STATISTICS ANA069127 AR746288 AA566223 BE090591 H93438 N73126 H93466 AA090928 AA095051 T29025 AW951071 L47277 L47277 G A1375913 BE384156 W24652 AA746288 AA566223 BE090591 H93033 N57027 AA504348 AA327653 AW959913 N53767 AA843715 AI453437 AW263710 AU76594 AA563483 AW873194 AW575166 AI128799 AI803319 AL042776 AW074313 AI887722 AI032284 AA447521 AI123885 N29334 AI354911 AW090687 AA236763 AA435535 AA236910 AA047124 AA236734 AW514610 H93467 AA962007 AI446783 AA127259 AI613495 AI686720 AI587374 AA936731 AA702453 AI859757
10			AA216786 Al251819 Al469227 AA806022 Al092324 N71868 AA968782 AA236919 AA809450 AA227220 AA765284 Al192007 AA766810 AA805794 AA729280 AA806238 AW768817 N71879 Al050686 AA505822 AA668974 Al688160 BE045915 AW466315 AA731314 AA649568 AA834316 AW591901 AW063876 AW294770 Al300266 Al336094 Al560380 AA721755 H09978 D20305 D29155 AW821790 BE150864 F01675
15	100780	458_127	BE561958 BE561728 BE397612 BE514391 BE269037 BE514207 BE562381 BE514256 BE514403 BE514250 BE397832 BE269598 BE559865 BE398831 BE560031 BE514199 BE560037 BE560454
	100830	4002_1	AC004770 W05005 AA356068 AA094281 H29358 T56781 AW875513 L37374 BE312466 BE311755 BE207105 BE293320 BE018115 AW239090 BE548830 AW247547 AA776062 BE397382 AA486713 T1011 T09340 AW498981 BE547280 AA356003 AW581520 AW875331 AA580720 AW375336 BE276873 BE408229 AW188148 BE255166 BE253761 AW793727 AW373141 AW581548 AA471223 AA305950 BE263976 AA626820 AW375336 BE276873 BE408229 AA09855 C00312 BE312741 BE407213 AA290352 AW298199 AW248553 AW297794 AW731722 BE300586 AW731972
20			AW615446 BE301599 AW615520 AA486714 AW440257 AA196516 AA564630 AA618079 AW192592 AW474985 AA604580 Al627461 AA765440 Al680394 AL135548 Al683224 Al581126 AW245096 AW194154 H29274 N70363 AA629758 AA580602 AA862006 Al863841 Al097667 Al928583 AL358774 BE243487 AA620553 AA653297 AA292690 T10110 Z38906 AA908544 AA340930 Al185438 T03328 T28844 Al687010 Al864965 AL972575 BE388740 T56780 AW373138 BF258717 AA699671
25	100906	4312_1	AU076916 BE298110 AW239395 AW672700 NM_003875 U10860 AW651755 BE297958 C03806 AI795876 AA644165 T36030 AW392852 AA446421 AW881866 AI469428 BE548103 T96204 R94457 N78225 AI564549 AW004984 AW780423 AW675448 AW087890 AA971454 AA305698 AA879433 AA535069 AI394371 AA928053 AI378367 N59764 AI364000 AI431285 T81090 AW674657 AW674987 AA897396 AW673412 BE063175 AW674408 AI202011 R00723 AI753769 AI460161 AW079585 AW275744 AI873729 D25791 BE537646 T81139 R00722
30	100930	16865_1	J04129 NM_002571 AA293088 AA477016 AA404631 T28299 AA476904 AA433965 AA430486 AA495907 AI151391 AA291495 AA402723 W25651 AA706816 AI826712 AW296294 AA293479 AI276581 AW044154 AI080180 AI417985 AI274168 AI474212 AA495908 AA635664 AI092114 AI804952 AA479874 AI597661 AI420511 AA479738 AA421417 AA421247 AA436220 AL047797 M34046 N42277 AA228076 W02698 AI420297 AA434011 AI369971 AA479731 AI865541 AI418020 AA421246 AA452764 AL048051
35	102221	3861_1	NM_006769 U24576 AW161961 AW160473 AW160465 AW160472 AW161069 AI824831 AW162635 AI990356 AW162477 AW162571 AI520836 AW162352 AW162351 AW162752 AI962216 AI537346 AA853902 H17667 BE045346 BE559802 BE255391 AA985217 AA235051 AI129757 AW366451 T34489 D56106 D56351 AI936579 AW023219 AW889332 AW889120 AW889322 AW889175 BE093702 AW889349 AA147646 AI952998 AA912579 AI143356 AW902211 R64717 AW157236 AI815242 D45274 AW263991 AA442920 AA129965 AL035713 AI923255 AI949082
40			A1142826 A1684160 A1701987 A1678954 A1827349 BE463635 AW628092 AW302281 AA493203 BE348856 BE536419 AW193969 AW673561 AW592609 A1224044 H43943 AA091912 R49632 R48353 A1568409 R48256 A1198046 H27986 H43899 A1678759 A1680310 A1624220 H17052 AA156410 N56062 A1699430 AA664529 T09406 T10459 AA627506 A1379584 N88633 AW022651 AA971281 AA248036 A1039197 A1914689 AA973825 AL047305 AA129866 A1798369 AW264348 A1445879 A165879 N67924 A1933507 A1216121 A1333174 T10972 A1375028 A1186756 A1273778 AA610487 A1797946 AA653903 AA903939 A1338587 A1278494 AW627595 AA904019
	101809	32963_1	M86849 AA315280 NM_004004 AA315269 BE142653 AA461400 AW802042 BE152893 AW383155 AA490688 AW117830 AW384563 AW384644 AW384566 AW378307 AW378323 AW839085 AA257102 AW378317 AW276060 AW271245 AW378298 AW384497 AI598114 AW264544 AI018136 AW021810 AA961504 AW066214 AW771489 AW192483 AI290266 AW192488 AW384490 AW007451 AW880895 AA554460 AA613715 AW200068 AJ783695 AI589498 AI917637 AW264471 AW384491 AI816732 AW368530 AW368521 AW368463 AA461087 AI341438 AI970613
45	102590	15932_1	A1040737 A1418400 AA947181 AA962716 A1280695 AW769275 AW023591 A1160977 AA055400 N71882 AA490466 AW243772 AW316636 AI076554 AW811702 N69323 H68912 AA257017 AI952506 H88913 AI91281 AA600714 BE465701 N64149 C00523 N64240 AA677120 R61573 BE005029 X98091 AA297307 BE537267 BE566138 BE566139 F11561 BE56750 BE568776 AW064005 BE566479 BE380035 BE567012 BE568634 BE566568 AA298060 BE566043 BE568813 BE568618 AA283070 BE566738 BE5667638 BE568667 BE566716 BE566433 U62136 AF649140 BE567057 BE567297 BE567403 BE568316 BE567400 BE566858 BA448772 AA071363 AW732642 BE564996
50			062136 AF-049140 BEB67/057 BEB67/297 BEB67/403 BEB687/30 BEB67/30
55			AU50943 AA661534 AA258061 AI090546 AA995167 AI051011 AA584421 AI026032 AW591338 AW589563 AA776914 AW024884 AA421002 F09219 BE464500 Al383595 AA954244 AA601583 AA737304 AA195549 AA805778 AI055876 AA164942 AW013961 AI672608 AW514211 D59441 AW582574 AA160935 BE566601 BE564612 BE566353 BE566195 BE566447 BE568302 BE568097 BE565470 BE564249 AL036217 AW749424 BE567494 AA102842 AA314761 AV661237 C14211 AA651866 AW798937 AA470605 AF112213 AL050318 T24804 AW28136 BE386341 BE263177 W16677 BE250224 BE563669 BE267405 BE546577 AV651354 AV651292
60	101977	29073_1	AT-172213 ALD90318 (24804 AWX-4813 BE3605431 BE260317 W10617 W10617 BE230224 BE30305 BE267 AVG BE3467 AVG BE34
65			AA553675 AI052791 AW059835 AI041906 AA814658 AW002059 AA729483 AI609301 AA994633 AA903651 AI459183 T95072 AW088630 AA128112 AI800091 AI561216 H17502 AW475072 AI819003 AI683272 AI262701 AW793140 T81787 R99586 AI275160 AI310420 AI698929 AA159174 AI827968 F30305 F30309 AA806662 AI091923 AW878722 AA583430 AW571913 AI674584 AA292533 AI079471 AA642325 AA719050 AW793172 AA305476 AW103745 T23459 N79525 AI784438 AA534551 AW193751 AI074360 BE281214 T32229 W25066 W01205 T63086 AW795348 AI361287 AW795353 AW795349 AA594759 AI400295 D11489 AI370689 AA482356 AA485295 W40151 AA564661 AW300745
70	102781	20812_1	ANY 59346 A1301267 AVY 59353 AV 59349 A3612604 A183409 AA996156 AW366963 AW366977 A1284860 AA846503 A1985064 AA844576 AA737921 AA873274 BE241546 BE241540 AA484058 AW468970 AA127876 AA159120 AW001568 AW795213 AW795258 AW795330 BE250589 BE387572 AA910895 AA161217 BE250380 W31500 T95167 A1719306 A1359224 BE258778 BE281230 BE410044 T33723 AW672694 AW410439 NM_006429 AF026292 T35505 BE542333 T08940 AU076737 AW247471
75	192/01	20012_1	BE393215 AW328640 BE542408 T32170 BE302544 T31955 BE206898 BE275738 T32570 BE386426 BE298746 BE389937 BE293991 BE315289 BE389578 R34739 R15312 BE279365 BE277756 AL036019 T33725 BE277779 BE302962 AL047294 BE276505 T09070 T33673 BE312580 AW387774 BE257175 AW674367 BE253331 BE270344 BE298931 BE273576 T32062 AI751831 BE618381 AA304899 BE252288 U46364 BE256790 BE207199 BE207199 BE251941 BE2507919 BE313955 BE269806 BE543623 BE279212 BE252289 T31699 BE262220 T31669 AA315781 AA192212 N84547 BE29737 BE259631 AA232179 A133144 T31292 AA315945 BE407301 BE251184 BE409006 AI880158 AI904003 AI904114 AW651768 AW651763 R58247 BE271897 U83843 C05298 BE261609 BE255973 AA351650 N84831 BE263537 AW452910 AA328465
80			A1904114 AW051/68 AW051763 K95247 BE211637 U63843 CU3248 BE261609 BE26357 AW315081 Ne46511 BE26357 AW32510 AW3
85			AW173930 AW311540 BESS5220 ANS65011 BESS 1104 Al600350 KW31353 AND65050 KW313643 AW615672 BES94607 AA483902 AW475032 BES78532 AW615636 AW732207 AW377638 AA321784 AA641629 AA633105 AA527640 AW129146 AW615672 BES94607 AA483902 AW475032 BES78532

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5		,	AAB72808 AI469388 AW105268 BE047301 AW591843 AW410066 AW517153 AI950495 AA746641 AI914878 AAB73185 AI696911 AA548625 AA911505 AA148762 AW674535 AI587329 BE328328 AW270348 AA158226 AW117705 AW474997 AW519183 AA614757 AW664383 AI082647 AW590973 AI476711 AA192213 N88741 BE464552 AW072679 AI453708 AA152166 AA805924 AI581078 AI125768 AW173484 AI961980 BE300766 AI199698 AI636792 AW247333 AW272861 AA078818 AA150012 AA551232 AA678821 AW873869 AW768266 AI660315 AA319210 AA814551 AA157994 AA318886 AI582962 AW089224 AI356098 AI343694 AW072598 N21054 AI301249 AA742924 H17917 AW328584
		:	AW248898 A1751830 AA907816 R08898 AW087989 A1828300 AA148596 A1269577 T33426 AA213571 A1973201 AA666279 R49612 A1573183 AW799762 AW410068 AW769666 A1962097 A1475204 D57490 AW517531 BE245270 AW474008 T33427 AW005731 A1795795 T23753 AW272981 T15747 AA552875 T23644 AW361289 A1758558 BE207435 AA876958 T03361 AA883569 F37533 AA582321 AW082524 R42212 AA973847 T18900 AA086202 A1559867 A1302418 AA948667 AA745670 T08939 T33724 T33722 BE621568 D57489 D25906 BE621151 F16510 C05966
10 15			T35127 AAG30427 AI933481 AA309426 AI918440 BE561854 BE618866 BE394675 BE296173 AW951687 BE383739 BE616141 BE312730 BE535351 AW080575 BE313330 BE616664 AI354390 AA847315 BE544509 BE515212 BE297833 BE278808 BE54484 AW090178 AI890664 BE546708 AW189943 BE274412 BE382399 BE266392 BE256949 BE280696 BE383237 BE261756 BE257721 BE312683 BE275476 BE514880 BE545314 BE313587 BE384537 BE386691 BE264813 AW592575 AI336332 AI278641 AI795791 BE222662 AW249316 AA314361 AL036012 AW402923 BE266845 AA075945 AA314436 BE384640 AW731769 AW957077 AA552234 AA573560 AW367038 AA313399 AI983873 BE410159 BE263803 BE514339 BE409073 BE281296 BE543396 BE395387 BE088360 BE546946 BE546570 BE390628 AA074638 AA301821 AW845230
	119221	102947_1	AW582379 AJ949222 AW029572 AA515843 AW272394 BE250234 C14322 W774050 AI074232 AA556284 BE048955 A148417 AI583145 AI473460 AI801688 AW573593 AI950741 AI628140 AW467921 R98105 A1149258 AI247584 AI078378 A1139850 AA489411 W24744 R98104 AI033826 AA699589 AI033120 N55544 W88984 AW970771 AA703362 AA099138 AA706792 AA046150 H98981 AI916674 AA953018 AI972749 AI921343 AA909044 AA094751 AI203124 AA582143 AI446654
20	125831 128192 113195	1522905_1 45743_3 178688_1	AW235415 R70377 AA099236 F20703 AA524436 R69484 H04043 D60988 D60337 A1204246 A1204250 A1194050 H83265 T63524 AA304359 AW960551 AI672874 A1749427 AA227777 AW027055 AA971834 T49644 T54122 AI983239 AI808233 T91264 T96544
25	119861	238266_1	A355945 A1709114 R72382 T48788 R48726 AW385418 A1095484 T49645 AA928653 AA570082 AW007545 T57178 AA516413 AA913118 T57112 AA564433 AA774503 AA367671 T59757 W78816 A1720806 A1633854 A1632086 A1668663 N70894 AW571809 A1383592 A1201348 W80715 N91880 AW963101 AA339011 AB033023 BE391906 BE275965 BE277872 BE003882 AA313774 BE019159 BE298024 BE299727 BE300011 BE390277 BE394764 N87550
30	112973		ABU33223 BE37930 BE279333 BE27712 BLGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGGG
35	129402 105936	47367_1 260931_1	MY2062 AF088057 W76255 AI827219 AI631461 AW449295 AI354957 AI913803 T62772 AI222040 T62921 T63781 AI678765 H12175 R14664 AI914049 AA995383 H08009 H19418 AW953728 AI358021 AA587361 AI269377 AA369905 AW957113 H27693 AI300474 H73776 W74397 AA578604 AI131018 W72331 AI719085 AA568348 AI859045 AI814819 AI888714 BE467470 AW131268 H19419 H27694 AI342165 AI914155 AA534872 BE018176 R60206 H11647 R45641 AI860466 BE301656 AI125453 AI498120 AA593735 AA879110 AI016404 T35018 AA588397 AW449767 AA470365 BE501139 AA588354 AI337500 AW078532 Z41279 AI125449 AA935725 AA404338
40	129466		L42593 NM_005554 L42601 BE183076 AJ541221 BE140567 L42610 V01516 J00269 AW275792 AW383052 AW380143 AJ541102 BE612846 AJ541344 AW238368 BE613405 BE615705 BE615530 BE615301 AW379823 AW794706 AA194806 AA194992 AW384024 AW384000 AA641239 AJ246504 AJ540333 AW238681 AA640939 AJ540863 AJ608860 AW862564 AW366725 AW366983 AW366870 AA596020 AW794721 AW794511
			AI591181 BE182523 AW794644 AW794620 AI935234 AI608903 AI608623 AW797060 AW084935 BE182517 BE182319 AI890082 AW238346 AW797012 BE182522 AW794838 AI608794 AW304289 AA147193 AA595995 AW381128 AW365720 AA583718 AI828416 BE122864 AW368343 AA431080 AW082039 AW380976 AA587144 AA443636 AW872937 AW794448 AW378382 AW085761 AW794718 AW263395 AA583587 AA583991 AA583994 AA586880 AA586880 AW368365 AI814460 AA586991 AI282829 AW378406 AA586721 AI609242 AA431973 AA232959
45		·	AIB31095 AW263854 AW378391 AW378415 AW378381 AA036990 AW238395 AI285446 BE208219 BE049526 AA583605 AA583918 AW366711 AI285580 AW082642 AI285712 AA582875 AW591216 AW368719 AW378408 BE122835 AA582976 BE350422 AA418228 AI541454 AI565930 AA583700 AA150575 AW238427 AI287474 AA912658 AA584223 AW238528 C17918 AW378169 AA159847 AI923797 AI609009 BE182479 AI915198 AW378114 AA147179 AA584239 AA150532 AW168862 AW085999 AW082480 AA659742 AW079703 AI872793 AA583981 AI824571 BE182316 BE182507 AA233331 AI824572 AI540586 D29492 BE182931 AA036948 BE551821 D29401 AW378365 C00141 D29181 D29567
50	100220 100355	45374_1 12538_1	AW103359 W95238 AI991663 AA587298 BE184608 AA099833 W95121 W95150 D29584 AI934111 D29456 D29533 AW265380 D29290 AW238463 AA121041 D29204 AA5959525 D29441 AW081840 AA587018 D29323 AA582891 BE182433 BE182437 BE158295 BE182434 AW015634 AA314369 AA290715 BE568683 AW629494 D28364 AW995678 AI907114 AA580734 AL041945 AA101515 AA121344 D78130 NM_003129 AA341650 T84166 AF098865 AA130976 BE089553
55		H05719 F13446	3 T66122 AW175590 F05344 A1114790 R12900 AA194871 AA132298 D78129 AA132213 AW948930 AW948919 AA263053 AW946593 AW946593 AM946840 AA278558 R50895 N26940 N40818 AW021265 AA054851 AA663379 AW948795 AW948893 AA400356 AW948911 N85024 W78844 AI341546 AI760182 AA286783 BEG17763 BEG17763 BEG17283 AW263690 BE049454 BEG17928 AW515038 AW950584 AA601009 AI079194 AA147204 AW083163 AA130981 AI218369 AA604794 AI806257 AI559556 AA232218 AA258085 AI471982 AA687949 AI143944 N30172 AA400196 AI769049 AI084342 AI221380 AA948469 AI802469 H05720 AA113270 AA158138 AA076231 AI559024 AI810962 AI133616 AA805106 AA101516 R40052 R50778 R43280 T66036 AW131924 AA114251 AA152331 F09650 AA580614 AA568927 C75491 Z38352 AA954959 C75606 W80742
60	100491 BE277805	34803_1 AA147951 AA600	D56165 M36981 X58965 NM_002512 BE379177 AA314836 BE256445 BE252016 AW248343 AI720933 AW085701 BE386050 BE619742 3113 BE253293 AI246588 AI183405 AI954174 AI126891 AI829101 AI123832 AW129670 AA471268 AW170242 AW873079 AA148011 AI608620 AA482961 AI003658 H43261 AA657978 AI735072 R83138 AA722002 AA626271 AW273877 BE464626 AA071483 AA429973 AA494342 AA620436 AA775597 AA775601 AA826847 AI192585 AA826359 AA411159 AI193419 AI204013 AA705323 AA716255 AI784611 AI081144 AI128227 AA828464 AI148911 AI493446 AI626084 AI189180 AI721196 AI190618 AA284987 AI128543 AA632064 AI333073 AI278470 AA131688
65			A1128227 AA828484 A114391 TAI93394 A1050084 A1109160 A121190 A1131016 A1204397 A112095 A204397 A12095 A204397 A12095 A1491768 AA937581 AA630055 AA934267 AN249841 AA583742 A1309756 AA961676 A1760860 AA557818 AA954238 H43655 A1302564 AA127545 A1609219 H20426 A1042292 A1056466 AA581836 W47002 AA422057 AA937673 F29757 AA829208 AW327462 AA372098 W02144 AA036805 AA487365 AA961037 A1139946 AA487250 AA737118 A1952504 A1242293 AA650552 A1708401 A1633133 AA630848 AA654317 F24128 A1434165 W46252 AW043879 A1033763 F37228 AA687890 N49087 AA876981 AA508474 A1914572 A1833284 F22253 AA026222 R50166 A1219267 N27095 AA495612 A1784222 A1289904 AA513146 AA528454 AA418700 F38721 A1880700 A1601170 A1862851 A1708633 AA524499
70			AA642220 AA496628 AI718709 W80579 AI720547 F20718 AA649943 AA688229 N40503 N46029 BE262669 BE391069 BE537538 AI510751 AI906968 AI318611 H46099 AI472604 T60667 AA373087 W32479 AA514034 BE619183 AA134672 AA127544 H26942 BE536689 AW327461 AA422139 AW262357 AW327348 F33510 AI630382 AW827126 F27133 AI335189 AW517589 W80471 AA885814 N89681 BE393173 AA617760 AA584268 AA460537 AA446261 H20425 N64040 AW276801 AA316367 AA071232 BE545409 AA308292 BE274447 AA380861 AA340038 AA341806 AA865579 AI018634 AI766314 AI919302 AA872367 AA991404 AI906981 AA88375 BE621012 AA505388 AA935192 AA290828
75	100518	13165_1	AGG 210 H50814 H44721 AW951723 AA514796 AA418708 AW673377 AA379622 AA977995 AA708224 AA708216 Al318249 Al318233 AA411160 AA026221 AA316774 AA486908 Al500094 AA096362 AW583742 BE536422 BE618653 R70203 AA131732 AA345048 BE562720 T28342 NM_004415 AL031058 M77830 BE149760 AW752599 AW848723 AW376697 AW376817 AW376699 AW848371 AW376782 AW848789 AW361413 AW849074 AW997139 AW799304 AW799309 BE077020 BE077017 BE185187 AW997196 BE156621 BE179915 BE006561 BE143155
80			AW890985 BE002107 AW103521 AA857316 AW383133 BE011378 AW170253 BE165750 AW886475 BE160433 J05211 BE082576 BE082584 BE004047 AW807238 AW377700 AW377699 BE082526 BE082505 BE082507 BE082514 AW178000 AW177933 AJ905935 AW747877 AW748114 BE148516 AW265328 AW847678 AW847678 AW365151 AW365151 AW365153 AW365156 AW365175 AW365175 AW365157 AW365154 AW066840 BE005272 AW365145 BE001925 BE182168 BE144243 BE001923 AJ951766 AJ434518 BE184920 BE184933 AJ284090 BE184941 AW804674 BE184924 C04715 W39488 AW995615 BE184948 BE159846 AW606653 AA099891 AA131128 AA337270 AA340777 AW384371 AA852212
85			R58704 AW366566 AW364859 AA025851 AA025852 AA455100 AA719958 AW352220 AW996245 BE165351 BE073467 AA377127 AW890264 AW609750 AW391912 AW849690 T87267 AW853812 AA852213 W74149 BE009090 AA056401 H91011 AW368529 AW390272 C18467

	wo	02/08644	PCT/US02/12476
5			AW674920 N57176 AA026480 AW576767 H93284 AA026863 AW177787 AA026654 AW177786 BE092134 BE092137 BE092136 AW177784 AI022862 BE091653 AW376811 AW848592 AA040018 BE185331 BE182164 AA368564 AW951576 T29918 AA131077 W95048 W25458 AW205789 H90899 N29754 W32490 R20904 BE167181 BE167165 N84767 H27408 H30146 Al190590 C03378 AI554403 AI205263 AA128470 AI392926 AF139065 AW370813 AW370827 AW798417 AW798780 AW798883 AW798569 R33557 AA149190 C03029 AW177783 AA088866 AW370829 AA247685 BE002273 AI760816 AI439101 AW879451 AI700963 AA451923 AI340326 AI590975 T48793 AI568096 AI142882 AA039975 AI470146 AA948936 BE067737 BE067786 W19287 AA644381 AA702424 AI417612 AI308554 AI686869 AI568892 AW190555 AI571075 AI220573 AA056527 AI471874 AI304772 AW517828 AI915596 AI627383 AI270345 AW021347 AW166807 AW105614 AI346078 AA552300 W95070
10			A4494069 AI911702 AA149191 AA026864 AI830049 AI887258 AW780435 AI910434 AIB19984 AI858282 AI078449 AI025932 AI860684 AI653678 AA026047 AA703232 D12062 AW192085 AA658154 AW514597 AW591892 T87181 AA782066 AW243B15 AW150038 AW268383 AW004633 AI927207 AA782109 AW473233 AI804485 AW169216 AI572669 AA602182 AW015480 AW771865 AI270027 AA961816 AA283207 AI076962 AI498487 AI348053 AI783914 H44405 AW799118 AA128330 AA515500 AA918281 W02156 AI905927 AA022701 W38382 R20795 T77861 AW860878
15	100528	45979_1	BE386801 AU077299 AA143755 BE302747 AA853375 U30162 BE274163 BE277479 BE408180 BE274874 C15000 AA047476 NZ7099 AI359165 AI538794 AI151283 AI863925 AW444977 AI207392 AA931283 AA443112 R40138 AW068538 AA351008 AA676972 R62503 AA916492 AW001865 H42334 H38280 AA121497 AA114137 AI750938 M17783 AA383786 BE274462 AI753182 C05975 AA347404 AW069298 AI754351 AI754044 AA188808 AA188879 AA565243 AL040655 AA456177 AI750722 AA045756 AA213580 C16936 AW578747 AW753731 H41692 N44761 R58560 R61260 AA039902 N59721 AW992543 R68380 AA149686 T29017 H03739 BE383822 BE387105 BE408251 BE410425 H41560 AA247591
20	100559	2260_1	BE389677 AI752233 AI568195 AA868004 AI424523 AW753720 AA852159 BE386803 NM_000094 L02870 D13694 S51236 M96984 AW946290 M66158 AI285422 D29523 AL119886 AW630655 L06862 AI884355 AW168737 T29085 AW797005 AW801340 AI355504 AW079048 AW801337 AI690455 AI972063 AW268565 W68588 AA587326 AA883498 AI033523 AW510356 AW591998 H98463 AL043852 AI150055 AI566239 AI624803 AA844717 H40670 AA922334 AI864424 AW615094 AW451233 AI302203 F31221 AI872170 W68589 AA904478 AI917631 AW014208 AW450759 AA847625 AI284033 AA848176 AA598507
25	100576 124357 101624 101625 135158	9986_1 genbank_N224 entrez_M55998 entrez_M57293 57963_1	B M55998

Tables 2A-8C were previously filed on November 9, 2001 in USSN 60/339,245 (18501-004100US)

Table 2A shows 504 genes down-regulated in lung tumors relative to normal lung and chronically diseased lung. Chronically diseased lung samples represent chronic non-malignant lung diseases such as fibrosis, emphysema, and bronchitis. These genes were selected from 59680 probesets on the Eos/Affymetrix Hu03 Genechip array. Gene expression data for each probeset obtained from this analysis was expressed as average intensity (AI), a normalized value reflecting the relative level of mRNA expression.

•	anpi coste		. prosecution	and well and one-just the only seeds as a re-	,	,,						•
	Pkey:	Unique	Eos probeset	identifier number								
	ExAccn:			number, Genbank accession number								
10	Unigenell		number									
10	Unigene	Title: Unigene	e gene tille 	or normal lung samples divided by the 80th percei	ntile of Al for s	innereir	nma and	SUISMOIR	s cell cam	dnoma lund	ı tumor	
	R1:	samples		or normal land samples divided by the cour percei	100 01 73 101 01	20110001011	ionia ana	oquamos	,		,	
	R2:	median	of Al for norm	at lung samples divided by 90th percentile of Al fo	r adenocarcino	oma and s	quamous	cell carcin	oma lung	tumor san	nples.	
	R3:	median	of Al for norm	al lung samples minus the 15th percentile of Al fo	r ali normal lun	a, chronic	ally disea	sed lung a	nd tumor	samples d	ivided by	
15		the 90th	percentile of	f Al for adenocarcinoma and squamous cell carcin	oma lung tumo	or samples	minus th	e 15th per	centile of	Al for all n	ormal	
		lung, ch	ronically disea	ased lung and tumor samples.					a humam			
	R4:	average	of Al for norm	nal lung samples divided by average Al for squam nal lung samples divided by the 90th percentile of	ous cell carcin	oma ano a rinomac	RIEHOCHIC	anoma iun	g tumors.	•		
	R5: R6:	nedian	of Al for norm	at lung samples divided by the 90th percentile of Al fo	r ell normal lur	a. chronic	ally disea	sed lung a	nd tumor	samples d	lvided by ti	he 90th
20	110.	oercent	le of Al for ad	enocarcinomas minus the 15th percentile of Al for	all normal lun	g, chronica	ally diseas	ed lung a	nd tumor	samples.		
	R7:	01/01/2010	of Al for norm	na) hung samples divided by the 90th nercentile of	Al for squamo	us cell car	cinomas.			•		
	R8:	median	of Al for norm	al lung samples minus the 15th percentile of Al fo	r all normal lur	ig, chronic	ally disea	sed lung a	ind tumor	samples d	ivided by ti	he 90th
		percent	ile of Al for sq	uamous cell carcinomas minus the 15th percentile	or Ar for all no	mai lung	, chronica	ily disease	io lung a	iu wiilui se	inhies.	
25	Pkey	ExAccn	UnigenelD	Unigene Title	R1	R2	R3	R4	R5	R6	R7	R8
			Jg.J	•								
	100095	Z97171	Hs.78454	myocilin; trabecular meshwork inducible	40.20							0.40
	100115	NM_002084	Hs.336920	glutathione peroxidase 3 (plasma).			2.30					3.46
30	100138	U83508	Hs.2463	angiopoietin 1		11.00	2.30					
30	100299 100306	D49493 U86749	Hs.2171 Hs.80598	growth differentiation factor 10 transcription elongation factor A (SII);		11.00				3.08		
	100308	NM_014767		KIAA0275 gene product								3.16
	100458	S74019	Hs.247979	Vpre-B	42.40							
·	100862	AA005247	Hs.285754	Hepatocyte Growth Factor Receptor						4.13		
35	100959	AA359129	Hs.118127	actin; alpha; cardiac muscle				125.60				
	101032	BE206854	Hs.46039	phosphoglycerate mutase 2 (muscle)	36.40			34.60				
	101081	AF047347	Hs.4880	armyloid beta (A4) precursor protein-bind solute carrier family 6 (neurotransmitte				193.20				
	101088 101125	X70697 AJ250562	Hs.553 Hs.82749	transmembrane 4 superfamily member 2				100.20		3.10		
40	101180	U11874	Hs.846	Interleukin 8 receptor; beta	•			54.86				
	101308	L41390		"Homo saplens core 2 beta-1,6-N-acetylgl	33.20							
	101330	L43821	Hs.80261	enhancer of filamentation 1 (cas-like do			0.00	36.40				
	101345	NM_005795	Hs.152175	Calcitonin receptor-like			2.29	70.55				
45	101346	A1738616 M26380	Hs.77348 Hs.180878	hydroxyprostaglandin dehydrogenase 15-(N lipoprotein lipase				70.55				3.54
73	101397 101414	NM_000066	Hs.38069	complement component 8; beta polypeptide							3.81	
	101435	NM_001100	Hs.1288	actin; alpha 1; skeletal muscle				34.60				
	101507	X16896	Hs.82112	interlaukin 1 receptor; type I	ı			37.60				
50	101530	M29874	Hs.1360	cytochrome P450; subfamily IIB (phenobar			0.54					4.25
50	101537	Al469059	Hs.184915	zinc finger protein; Y-linked		5.50	2.54					
	101542 101545	NM_000102 BE246154	Hs.1363 Hs.154210	cytochrome P450; subfamily XVII (steroid EDG1; endothelial differentiation, sphin	39.40	9.30						
	101554	BE207611	Hs.123078	thyroid stimulating hormone receptor		13.00			•			
	101560	AW958272	Hs.83733	Intercellular adhesion molecula 2, exon								3.38
55	101574	M34182	Hs.158029	protein kinase; cAMP-dependent; catalyti						4.37		0.00
	101605	M37984	Hs.118845	troponin C; slow	00.00							3.80
	101621	BE391804	Hs.62661	guanylate binding protein 1; interferon-	30.20						2.75	
	101680 101829	AA299330 AW452398	Hs.1042 Hs.129763	Sjogren syndrome antigen A1 (52kD; ribon solute carrier family 8 (sodium/calcium						3.37		
60	101842	M93221	Hs.75182	mannose receptor; C type 1				38.20				
	101961	AW004056	Hs.168357	"Hs-TBX2=T-box gene (T-box region) [huma			2.32					
	101994	T92248	Hs.2240	uteroglobin			0.45					6.85
	102020	AU077315	Hs.154970	transcription factor CP2			2.45					6.75
65	102091 102112	BE280901 AW025430	Hs.83155 Hs.155591	aldehyde dehydrogenase 7 forkhead box F1	54.60							0.70
05	102112	AA723157	Hs.73769	folate receptor 1 (adult)	01100							3.98
	102202	NM_000507	Hs.574	fructose-bisphosphatase 1								3.62
	102241	NM_007351		Multimerin			2.32					
70	102310	U33839		Accession not listed in Genbank	00.40	7.00						
70	102397	U41898	Un aannen	"Human sodium cotransporter RKST1 mRNA, "Homo sapiens skeletal muscle LIM-protel	29.40							3.75
	102571 102620	U60115 AA976427	Hs.239069 Hs.121513	Human clone W2-6 mRNA from chromosome X						3.07		00
	102636	U67092	110.121010	"Human ataxia-telanglectasia locus prote			2.40					
	102667	U70867	Hs.83974	solute carrier family 21 (prostaglandin			3.15					
75	102675	U72512	Hs.7771	"Human B-cell receptor associated protei						3.56		A 54
	102698	M18667	Hs.1867	progastricsin (pepsinogen C)					12.00			4.51
	102727	U79251	Hs.99902	opioid-binding protein/cell adhesion mol conficotropin releasing hormone	37.40				12.00			
	102852 103026	V00571 X54162	Hs.75294 Hs.79386	thyroid and eye muscle autoantigen D1 (6	UP. 10				13.00			
80	103028	X54380	Hs.74094	pregnancy-zone protein	28.80							
	103098	M86361		Human mRNA for T cell receptor, clone IG		_			10.00			
	103117	X63578	Hs.295449	parvalbumin		6.00	0.4-					
	103241	X76223	70-00	H.saplens MAL gene exon 4			2.47 2.69					
85	103280	U84722 Y16791	Hs.76206	Cadherin 5, VE-cadherin (vascular epithe keratin; hair; acidic; 5			2.03				2.16	
65	103360	1 10131	Hs.73082	National, Itali, deletic, J								

	w	O 02/08	5443							PCT/	US02/1	12476
	103496	Y09267	Hs.132821	flavin containing monooxygenase 2								5.97
	103508	Y10141 NM 001843	Un 142424	"H.sapiens DAT1 gene, partial, VNTR" contactin 1			2.40			3.27		
	103561 103569	NM_005512		glycoprotein A repetitions predominant			2.99					
5	103575	Z26256		"H.sapiens isoform 1 gene for L-type cal						4.18 3.44		
	103627 103767	Z48513 BE244667	Hs.296155	H.sapiens XG mRNA (clone PEP6) CGI-100 protein						3.44	2.25	
	103767	AA187101	Hs.213194	Hypothetical protein MGC10895; sim to SR				46.55				
10	104078	AA402801	Hs.303276	EŜTs						3.05 3.54		
10	104326 104352	AW732858 BE219898	Hs.143067 Hs.173135	ESTs dual-specificity tyrosine-(Y)-phosphoryl						3.16		
	104398	AI423930	Hs.36790	ESTs; Weakly similar to putative p150 [H	64.80							3.38
	104473 104493	AI904823 AW960427	Hs.31297 Hs.79059	ESTs ESTs; Moderately similar to TGF-BETA REC			2.47					3.30
15	104495	AW975687	Hs.292979	ESTs	28.60							
	104595	A1799603	Hs.271568	ESTs		6.00				3.42		
	104597 104659	A1364504 AW969769	Hs.93967 Hs.105201	ESTs; Weakly similar to Slit-1 protein [ESTs	34.00	0.00						
	104686	AA010539	Hs.18912	ESTs		11.00						
20	104691	U29690	Hs.37744	ESTs; Beta-1-adrenergic receptor	56.80			60.40				
	104764 104776	A1039243 AA026349	Hs.278585	ESTs ESTs	34.20			•••••				
	104825	AA035613	Hs.141883	ESTs	44.00		3.03	•				
25	104865 104942	T79340 NM_016348	Hs.22575 Hs.10235	Homo sapiens cDNA: FLJ21042 fis, clone C ESTs	41.20							3.27
2-0	104989	R65998	Hs.285243	ESTs				40.00			•	2 20
	105062	AW954355	Hs.36529 Hs.38163	ESTs	34.20							3.20
	105101 105173	H63202 U54617	Hs.8364	ESTS ·	34.20							4.17
30	105194	R06780	Hs.19800	ESTs		16.00	2 24					
	105226 105256	R58958 AA430650	Hs.26608 Hs.16529	ESTs transmembrane 4 superfamily member (tetr			2.34 2.72					
	105394	BE245812	Hs.8941	ESTs			2.61					
35	105647	Y09306	Hs.30148 Hs.18142	homeodomain-interacting protein kinase 3 arrestin; beta 2	33.60							3.59
33	105789 105817	AF106941 AA397825	NS. 10 142	synaptopodin						4.46		
	105847	AW964490	Hs.32241	ESTs			3.43	35.40				
	105894 105999	AI904740 BE268786	Hs.25691 Hs.21543	calcitonin receptor-like receptor activi ESTs		7.00	3.43					
40	106075	AA045290	Hs.25930	ESTs				42.60				
	106178 106381	AL049935 AB040916	Hs.301763 Hs.24106	KIAA0554 protein ESTs	34.80				12.00		•	
	106467	AA450040	Hs.154162	ADP-ribosylation factor-like 2	•				,	3.69		
45	106536	AA329648	Hs.23804	ESTs				96.40 47.20		•		
43	106569 106605	R20909 AW772298	Hs.300741 Hs.21103	sorcin Homo saplens mRNA; cDNA DKFZp564B076 (fr				220.40				
	106842	AF124251	Hs.26054	novel SH2-containing protein 3	00.00		2.55					
•	106844 106870	AA485055 Al983730	Hs.158213 Hs.26530	sperm associated antigen 6 serum deprivation response (phosphatidy)	39.20		2.28					
50	106943	AW888222	Hs.9973	ESTs								4.28
	106954	AF128847	Hs.204038	ESTs					10.45		·	4.32
	107106 107163	AA862496 AF233588	Hs.28482 Hs.27018	ESTs ESTs			2.57					
EE	107201	D20378	Hs.30731	EST		8.00				3.84		
55	107238 107376	D59362 U90545	Hs.330777 Hs.327179	EST solute carrier family 17 (sodium phospha		10.67						
	107530	Y13622	Hs.85087	latent transforming growth factor beta b			2.32	04.00				
	107688 107706	AW082221 AA015579	Hs.60536 Hs.29276	ESTs ESTs	28.40			34.60				
60	107723	AA015967	113.23210	EST	20.10					3.29		
	107727	AA149707	Hs.173091	DKFZP434K151 protein				80.80 51.40				
	107750 107751	AA017291 AA017301	Hs.60781 Hs.235390	ESTs ESTs				01.40		3.14		
65	107873	AK000520	Hs.143811	ESTs		9.00				3.65		
65	107899 107994	BE019261 AA036811	Hs.83869 Hs.48469	ESTs; Weakly similar to IIII ALU SUBFAMI ESTs				44.60		0.00		
	107997	AL049176	Hs.82223	Human DNA sequence from clone 141H5 on c				32.00				
	108041	AW204712 AI797341	Hs.61957 Hs.165195	ESTs ESTs				30.80			4.75	
70	108048 108338	AA070773	па. 103133	"zm53g11.s1 Stratagene fibroblast (#9372			2.33					
	108434	AA078899		"zm94b1.s1 Stratagene colon HT29 (#93722						3.06	2.92	
	108447 108480	AA079126 AL133092	Hs.68055	"zm92a11.s1 Stratagene ovarian cancer (# ESTs				34.00		0.00		
7-	108499	AA083103		"zn1b12.s1 Stratagene hNT neuron (#93723					10.00			3.36
75	108535	R13949	Hs.226440	Homo sapiens clone 24881 mRNA sequence "zn11f6.s1 Stratagene hNT neuron (#93723					19.00 12.00			
	108550 108604	AA084867 AA934589	Hs.49696	ESTs			2,33					
•	108625	AW972330	Hs.283022	ESTs							3.42	5.82
80	108629 108655	AA102425 AA099960		"zn24c6.s1 Stratagene neuroepithelium NT "zm65c6.s1 Stratagene fibroblast (#93721		7.00					U.TE	
-0	108756	AA127221	Hs.117037	Homo saplens mRNA; cDNA DKFZp564N1164 (f	00.00	6.05						
	108864	Al733852 Al.138272	Hs.199957 Hs.62713	ESTs ESTs	28.80 32.80							
• -	108895 108921	AL138272 Al568801	Hs.71721	ESTS				57.80				
85	108967	AA142989	Hs.71730	ESTs	28.80							

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	109001	AI056548	Hs.72116	ESTs, Moderately similar to hedgehog-int			2.57				0.44	
	109003	AA147497	Hs.71825	ESTs EST		5.60					2.11	
	109004 109065	AA156235 AA161125	Hs.139077 Hs.252739	EST		3.00			10.00			
5	109250	H83784	Hs.62113	ESTs; Weakly similar to PHOSPHATIDYLETHA							3.44	
	109490	AA233416	Hs.139202	ESTs			0.40				2.92	
	109510 109578	A1798863 F02208	Hs.87191 Hs.27214	ESTs ESTs		10.00	2.40					
	109601	F02695	Hs.311662	EST		10.00		40.80				
10	109613	H47315	Hs.27519	ESTs				54.40				
	109650	R31770	Hs.23540	ESTs	31.20	8.40						
	109682 109724	H18017 D59899	Hs.22869 Hs.127842	ESTs ESTs		0.40		29.40				
	109782	AB020644	Hs.14945	long fatty acyl-CoA synthetase 2 gene					8.00			
15	109833	R79864	Hs.29889	ESTs		10.00	C 40					
	109837	H00656	Hs.29792 Hs.282982	ESTs ESTs			6.49				2.75	
	109977 109984	T64183 Al796320	Hs.10299	ESTs				107.00				
	110146	H41324	Hs.31581	ESTs; Moderately similar to SYNTAXIN 18							2.22	
20	110271	H28985	Hs.31330	ESTs	44.00					3.48		
	110280 110420	AW874263 R93141	Hs.32468 Hs.184261	ESTs ESTs	44.20			32.00				
	110578	T62507	Hs.11038	ESTs	28.40							
0.5	110634	R98905	Hs.35992	ESTs					20.00			4.40
25	110726	AW961818	Hs.24379	potassium voltage-gated channel; shaker-				56.80				4.15
	110837 110875	H03109 N35070	Hs.108920 Hs.26401	ESTs; Weakly similar to semaphorin F [H. tumor necrosis factor (ligand) superfami			3.13	00.00				
	110894	R92356	Hs.66881	ESTs; Moderately similar to cytoplasmic		5.33						
20	110971	A1760098	Hs.21411	ESTs	00.40			44.60				
30	111023 111057	AV655386 T79639	Hs.7645 Hs.14629	ESTs ESTs	32.40				17.14			
	111247	AW058350	Hs.16762	Homo saplens mRNA; cDNA DKFZp564B2062 (f							4.58	
	111330	BE247767	Hs.18166	KIAA0870 protein								3.42
35	111374	BE250726	Hs.283724	ESTs; Moderately similar to HYA22 [H.sap				33.20				3.91
33	111442 111737	AW449573 H04607	Hs.181003 Hs.9218	ESTs ESTs				53.00				
	111747	AI741471	Hs.23666	ESTs	46.20							
	111807	R33508	Hs.18827	ESTs		16.00				2.01		
40	111862 112045	R37472 A1372588	Hs.21559 Hs.8022	EST TU3A protein						3.91	2.74	
40	112045	R43713	Hs.22945	EST							4.92	
	112214	AW148652	Hs.167398	ESTs					13.00			
	112263	R52393	Hs.25917	ESTs		9.00	2.43					
45 ·	112314 112324	AW206093 R55965	Hs.748 Hs.26479	ESTs limbic system-associated membrane protei		5.00			14.00			
	112362	AW300887	Hs.26638	ESTs; Weakly similar to CD20 receptor [H			2.49					
	112380	H63010	Hs.5740	ESTs		0.00	2.34					
	112425 112473	AA324998 R65993	Hs.321677 Hs.279798	ESTs; Weakly similar to IIII ALU SUBFAMI pregnancy specific beta-1-glycoprotein 9		8.00				4.53		
50	112492	N51620	Hs.28694	ESTs				29.80				
	112541	AF038392	Hs.116674	ESTs			0.07			3.62		
	112620 112623	R80552 AW373104	Hs.29040 Hs.25094	ESTs ESTs			2.37 2.26					
	112867	T03254	Hs.167393	ESTs					12.00			
55	112894	T08188	Hs.3770	ESTs		6.50						
	112954	AA928953	Hs.6655	ESTS		7.00						4.39
	113029 113086	AW081710 AA346839	Hs.7369 Hs.209100	ESTs; Weakly similar to IIII ALU SUBFAMI DKFZP434C171 protein								4.47
	113140	T50405	Hs.175967	ESTs					10.00			•
60	113252	NM_004469		c-fos Induced growth factor (vascular en		14.00				0.70		
	113257	AI821378	Hs.159367	ESTs ESTs						3.72 3.60		
	113394 113437	T81473 T85349	Hs.177894 Hs.15923	EST	35.00					0.00		
	113454	Al022166	Hs.16188	ESTs		6.00						
65	113502	T89130	V- 40000	ESTs	39.60							3.88
	113552 113645	AI654223 T95358	Hs.16026 Hs.333181	ESTs ESTs							2.58	0.00
	113691	T96935	Hs.17932	EST				38.20				
70	113706	AA004693	Hs.269192	ESTs						3.09		
70	113883	U89281	Hs.11958	oxidative 3 alpha hydroxysteroid dehydro Homo saglens mRNA; cDNA DXFZp586B0220 (f	30.40		2.31					
	113924 114035	BE178285 W92798	Hs.170056 Hs.269181	ESTs	30.40				13.00		•	
	114058	AK002016	Hs.114727	ESTs								5.00
75	114084	AA708035	Hs.12248	ESTs			2 24	40.60				
75	114121 114124	H05785 W57554	Hs.25425 Hs.125019	ESTs Human lymphoid nuclear protein (LAF-4)		7.00	2.31					
	114275	AW515443	Hs.306117	Interleukin 13 receptor; alpha 1		6.00						
	114297	AA149707	Hs.173091	DKFZP434K151 protein				48.80		0.45		
80	114427	AA017176	Hs.33532	ESTs; Highly similar to Miz-1 protein [H					10.00	3.45		
οU	114449 114452	AA020736 Al369275	Hs.243010	"ze63b11.s1 Soares retina N2b4HR Homo sa ESTs, Moderately similar to RTC0_HUMAN G		14.00			10.00			
	114609	AA079505		"zm97a5.s1 Stratagene colon HT29 (#93722						3.13		
	114648	AA101056	11. 4550-:	"zn25b3.s1 Stratagene neuroepithelium NT				35.40				2 42
85	114731 114762	BE094291 AA146979	Hs.155651 Hs.288464	Homo sapiens HNF-3beta mRNA for hepatocy ESTs	33.00							3.42
5 5	114702	POLITORIS	10.200104	LUIU	55.50							

	W	O 02/086	5443							PCT/	US02/1	2476
	114776 115009	AA151719 AA251561	Hs.95834 Hs.48689	ESTs ESTs	34.40 30.20							
	115272	AW015947	115.40003	ESTs; Weakly similar to hypothetical L1	32.60							
- 5	115279	AW964897	Hs.290825	ESTs ESTs		6.00			12.00			
3	115302 115365	AL109719 AW976252	Hs.47578 Hs.268391	ESTs					12.00	3.32		
	115559	AL079707	Hs.207443	ESTs				48.00 56.20				
	115566 115683	A1142336 AF255910	Hs.43977 Hs.54650	ESTs ESTs, Weakly similar to (defline not ava	31.40							
10	115744	AA418538	Hs.43945	ESTs; Highly similar to dJ1178H5.3 [H.sa				33.60 74.40				
	115819 115949	AA486620 AI478427	Hs.41135 Hs.43125	Endomucin 2 ESTs			3.18	14.40				
	115965	AA001732	Hs.173233	ESTS				388.80 33.20				
15	116035 116049	AA621405 AA454033	Hs.184664 Hs.41644	ESTs . ESTs				45.80				
	116081	Al190071	Hs.55278	ESTs			3.06			3.57		
	116082 116213	AB029498 AA292105	Hs.59729 Hs.326740	ESTs leucine rich repeat (in FLII) interactin	50.60		3.00					
20	116228	Al767947	Hs.50841	ESTs; Weakly similar to tuftelin [M.musc		c 00	3.85					
20	116250 116419	N76712 Al613480	Hs.44829 Hs.47152	ESTs ESTs; Weakly similar to testicular tekti		6.00		30.00				
	116617	D80761	Hs.45220	EST	47.00	•	2.27					
	116784 116835	AB007979 N39230	Hs.301281 Hs.38218	tenascin R (restrictin; Janusin) ESTs	47.20	•		41.20				
25	116970	AB023179	Hs.9059	KIAA0962 protein				04.00	11.00			
	117023 117027	AW070211 AW085208	Hs.102415 Hs.130093	ESTs ESTs	49.40			91.00				
	117036	H88908	Hs.41192	EST		0.67		32.60				
30	117110 117209	AA160079 W03011	Hs.172932 Hs.306881	ESTs .		8.67		30.60				
50	117325	N23599	Hs.43396	ESTs					9.29	2 40		
	117454 117475	N29569 N30205	Hs.44055 Hs.93740	ESTs	44.00					3.19		
25	117543	BE219453	Hs.42722	ESTs		16.00			40.00			
35	117567 117570	AW444761 N48649	Hs.44565 Hs.44583	ESTs ESTs					12.00 11.00			
	117600	N34963	Hs.44676	EST		0.00				3.74		
	117730 117791	N45513 N48325	Hs.46608 Hs.93956	ESTs EST		6.00 9.00						
40	117929	N51075	Hs.47191	ESTs		0.00		29.20				
	117990 118224	AA446167 N62275	Hs.47385 Hs.48503	ESTs EST	31.40	8.00						
	118244	N62516	Hs.48556	ESTs	32.80		0.40					
45	118357 118446	AL109667 N66361	Hs.124154 Hs.269121	Homo sapiens mRNA full length Insert cDN ESTs	•		2.40 2.28					
	118447	N66399	Hs.49193	EST	30.80					3.10		
	118530 118549	N67900 N68163	Hs.118446 Hs.322954	ESTs EST						3.41		
50	118823	W03754	Hs.50813	ESTs; Weakly similar to long chain fatty			3.94			3.58		
50	118862 118935	W17065 Al979247	Hs.54522 Hs.247043	ESTs KIAA0525 protein				33.00		3.36		
	118944	Al734233	Hs.226142	ESTs; Weakly similar to !!!! ALU SUBFAMI		44.00			11.43			
	118995 119073	N94591 BE245360	Hs.323056 Hs.279477	ESTs ERG-2/ERG-1; V-ets avian erythroblastosi		14.00		52.60				
55	119268	T16335	Hs.65325	EST	31.40					3.50		
	119514 119824	W37937 W74536	Hs.184	Accession not listed in Genbank advanced glycosylation end product-speci			2.75					
	119831	AL117664	Hs.58419	DKFZP586L2024 protein				33.80				3.21
60	119861 119889	W78816 W84346	Hs.49943 Hs.58671	ESTs; Moderately similar to IIII ALU SUB ESTs				30.03				
	119921	W86192	Hs.58815	ESTs	29.00					3.80		
	120082 120094	H80286 AAB11339	Hs.40111 Hs.124049	ESTs ESTs		6.00				5.00		
65	120132	W57554	Hs.125019	Human lymphoid nuclear protein (LAF-4)		12.00		36.60				
65	120378 120404	AA223249 AB023230	Hs.285728 Hs.96427	ESTs KIAA1013 protein	39.40	12.00						
	120504	AA256837		ESTs	22.00				8.00			
	120512 120667	N55761 AA287740	Hs.194718 Hs.78335	ESTs microtubule-associated protein; RP/EB fa	33.00							4.18
70	120777	AA287702	Hs.10031	KIAA0955 protein				46.60 39.00				
	121082 121191	AA398722 AA400205	Hs.104447	ESTs ESTs	41.60			33.00				
	121248	AA400914	Hs.97827	EST					12.00		5.08	
75	121363 121366	A1287280 A1743515	Hs.97933	ESTs ESTs					20.00			
-	121483	A1660332	Hs.25274	ESTs; Moderately similar to putative sev				30.20		3.32		
	121518 121545	AA412155 AA412442	Hs.98132	ESTs ESTs			2.29	55.20				
80	121622	AA416931	Hs.126065	ESTs		9.00		34.80				
90	121665 121709	AA416556 Al338247	Hs.98234 Hs.98314	ESTs Homo sapiens mRNA; cDNA DKFZp586L0120 (f	34.80			J-1.00				
	121730	Al140683	Hs.98328	ESTs	38.80	7.00						
	121740 121772	AA421138 Al590770	Hs.98334 Hs.110347	EST Homo sapiens mRNA for alpha integrin bin	36.20	7.00						
85	121821	AL040235	Hs.3346	ESTs								3.61

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	121835	AB033030	Hs.300670	ESTs .			2.34					
	121841	AA427794	Hs.104864	ESTs			2.61				2.25	
	121885 121888	AA934883 AA426429	Hs.98467 Hs.98463	ESTs ESTs							2.92	
5	121938	AA428659	Hs.98610	ESTs				46.80				
-	121950	AA429515		EST				31.40				
	122030	AA431310	Hs.98724	ESTs	34.40						2 50	
	122054	AA431725	Hs.98746	EST ESTs; Moderately similar to bithoraxoid-	49.40						3.58	
10	122211 122233	AA300900 AA436455	Hs.98849 Hs.98872	EST	29.80							
	122247	AA436676	Hs.98890	EST				39.80				
	122253	AA436703	Hs.104936	ESTs; Weakly similar to hypothetical pro		9.00				2.00		
	122266 122285	AA436840 AA436981	Hs.98907 Hs.121602	EST EST						3.60 3.14		
15	122409	AA446830	Hs.99081	ESTs	30.80					0.14		
	122485	AA524547	Hs.160318	phospholemman			2.65					
	122697	AA420683	Hs.98321	Homo saplens cDNA FLJ14103 fis, clone MA		15.00						
	122772 122831	AW117452 AI857570	Hs.99489 Hs.5120	ESTs ESTs		6.67				3.37		
20	122913	A1638774	Hs.105328	ESTs				32.20		0.0.		
	123049	BE047680	Hs.211869	ESTs				41.80				
	123076	Al345569	Hs.190046	ESTs	35.80						2.58	
	123136 123309	AW451999 N52937	Hs.194024 Hs.102679	ESTs ESTs					19.00		2.30	
25	123455	AA353113	Hs.112497	ESTs				82.80	10.00			
	123691	AA609579	Hs.112724	ESTs						3.95		
	123756	AA609971	Hs.112795	EST	35.40							
	123802 123837	AA620448 Ai807243	Hs.112893	Homo sapiens clone 24760 mRNA sequence ESTs	58.00			32.40				
30	123844	AA938905	Hs.120017	olfactory receptor; family 7; subfamily			2.63	02.70				
	123936	NM_004673	Hs.241519	ESTs	29.00							
	123987	C21171	Hs.95497	ESTs; Weakly similar to GLUCOSE TRANSPOR	00.40			70.60				
	124013	Al521936 R40290	Hs.107149 Hs.124685	ESTs; Weakly similar to PTB-ASSOCIATED S ESTs	28.40				13.00			
35	124160 124205	H77570	Hs.108135	ESTs					10.00	4.74		
	124226	AA618527	Hs.190266	ESTs			2.35					
	124246	H67680	Hs.270962	ESTs		17.00		29.40				
	124348 124358	Al796320 AW070211	Hs.10299 Hs.102415	ESTs *yw35g11.s1 Morton Fetal Cochlea Homo sa		17.00	3.07					
40	124409	Al814166	Hs.107197	ESTs			5.51			3.14		
	124442	AW663632	Hs.285625	TATA box binding protein (TBP)-associate			2.48					
	124468	N51413	Hs.109284	ESTS				30.80				6.03
	124479 124519	AB011130 Al670056	Hs.127436 Hs.137274	calcium channel; voltage-dependent; alph ESTs; Weakly similar to SPLICEOSOME ASSO			2.50					0.00
45	124711	NM_004657		serum deprivation response (phosphatidy)	59.20							
	124866	AJ768289	Hs.304389	ESTs		8.00		07.00				
	124874 125097	BE550182 AW576389	Hs.127826 Hs.335774	ESTs ESTs				37.60	10.00			
	125179	AW206468	Hs.103118	ESTs					10.00	3.12		
50	125200	AW836591	Hs.103156	ESTs							2.79	
	125299	T32982	Hs.102720	ESTs	29.00			34.20				
	125400 125810	AL110151 H00083	Hs.128797	DKFZP586D0824 protein aryl hydrocarbon receptor-interacting pr	32.20							
	126176	BE242256	Hs.2441	KIAA0022 gene product		12.00	•					
55	126303	D78841		HUM525A05B Human placenta polyA+ (TFuji				33.60				
	126403	AW629054	Hs.125976	ESTs; Weakly similar to metalloprotease/ ESTs; Weakly similar to HC1 ORF [M.muscu	35.80			29.80				
	126507 126773	AL040137 AA648284	Hs.23964 Hs.187584	ESTs	39.60			25.00				
	127307	AW962712	Hs.126712	ESTs: Weakly similar to pIL2 hypothetica	28.80							
60	127462	AA760776	Hs.293977			9.00		34.40				
	127486 127572	AW002846 AA594027	Hs.105468 Hs.191788	ESTs ·		5.00	2.36					
	127609	X80031	Hs.530	ESTs				29.40				
CE	127832	AW976035	Hs.292396	ESTs				37.20			4.40	
65	127898	AA774725	Hs.128970 Hs.125983	ESTs ESTs				38.40			4.42	
	128073 128101	AW340720 AA905730	Hs.128254	ESTs		7.33		00.40				
	128149	NM_012214	Hs.177576	mannosyl (alpha-1;3-)-glycoprotein beta-							2.58	
70	128212	W27411	Hs.336920	glutathione peroxidase 3 (plasma)			3.09	34.40				
70	128333 128364	W68800 N76462	Hs.12126 Hs.269152	ESTs; Weakly similar to LR8 [H.sapiens] ESTs; Weakly similar to ZINC FINGER PROT		10.00		34.40				
	128426	Al265784	Hs.145197	ESTs							4.31	
	128598	AA305407	Hs.102308	potassium inwardly-rectifying channel; s	31.20							
75	128634	AA464918	II- 00707	ESTs; Moderately similar to IIII ALU SUB				41.60 87.00				
13	128687 128726	AW271273 Al311238	Hs.23767 Hs.104476	ESTs ESTs				07.00				4.02
	128773	NM_004131	Hs.1051	granzyme B (granzyme 2; cytotoxic T-lymp					9.00			
	128833	W26667	Hs.184581	ESTs			2.00					3.76
80	128870 128878	H39537 R25513	Hs.75309 Hs.10683	eukaryotic translation elongation factor ESTs			2.66			3.10		
50	128885	AF134803	Hs.180141	cofilin 2 (muscle)					11.00	UV		
	128998	W04245	Hs.107761	ESTs; Weakly similar to PUTATIVE RHO/RAC							3.21	
	129000	AA744902	Hs.107767	ESTs; Moderately similar to CaM-KII inhi						3.17		3.68
85	129038 129098	AW156903 AW580945	Hs.108124 Hs.330466	ribosomal protein L41 ESTs	34.60					0.17		
	123030		, 10.000700									

129210	4.09 4.05 4.09
129262 BE222198 Hs.109843 ESTs 3.30 129301 AF182277 Hs.330780 Human cytochrome P450-IIB (hitB3) mRNA; 129331 AW16766B Hs.279772 ESTs; Highly slmilar to CGI-38 protein [
129301 AF182277 Hs.330780 Human cytochrome P450-IIB (hIB3) mRNA; 129331 AW167668 Hs.279772 ESTs; Highly similar to CGI-38 protein [129381 AW245805 Hs.10903 claudin 5 (transmembrane protein deleted vasoactive intestinal peptide receptor 1 129595 U09550 Hs.1154 oviductal glycoprotein 1; 120kD 10.00 129613 AW978517 Hs.172847 ESTs; Weakly similar to collagen globa 1 3.40	
129381 AW245805 Hs.110903 claudin 5 (transmembrane protein deleted 2.93 129565 X77777 Hs.198726 vasoactive intestinal peptide receptor 1 160.80 129595 U09550 Hs.1154 oviductal glycoprotein 1; 120kD 10.00 129613 AW978517 Hs.172847 ESTs: Weakly similar to collagen alpha 1 3.40	4.09
129565 X77777 Hs. 198726 vasoactive intestinal peptide receptor 1 160.80 129595 U09550 Hs. 1154 oviductal glycoprotein 1; 120kD 10.00 129613 AW978517 Hs. 172847 ESTs: Weakly similar to collagen alpha 1 3.40	
129613 AW978517 Hs.172847 ESTs: Weakly similar to collegen alpha 1 3.40	
10 129782 AW016932 Hs.104105 EST 9.00	
129950 F07783 Hs.1369 decay accelerating factor for complement 87.80	
129958 R27496 Hs.1378 annexin A3 44.60 129959 AL036554 Hs.274463 defensin; alpha 1; myeloid-related seque 2.72	
130160 A3305688 Hs.267695 UDP-Gal:betaGlcNAc beta 1:3-galactosyltr 42.20	
15 130259 NM_000328 Hs.153614 retinitis plgmentosa GTPase regulator 2.54	
130273 AW972422 Hs.153863 MAD (mothers against decapentaplegic; Dr 51.60 130312 AF056195 Hs.15430 DKFZP586G1219 protein 3.16	
130436 NM_001928 Hs.155597 D component of complement (adipsin)	4.11
130523 AA999702 Hs.214507 ESTs 4.77 20 130799 AB028945 Hs.12696 ESTs 6.00	
20 130799 AB028945 Hs.12696 ESTs 5.00 130885 NM_005883 Hs.20912 adenomatous polyposis coli like 3.54	
131002 AL050295 Hs.22039 KIAA0758 protein	3.50
131012 AL039940 Hs.202949 KIAA1102 protein 20.00 131031 NM_001650 Hs.288650 aquaporin 4 41.20	
25 131061 N64328 Hs. 268744 ESTs; Moderately similar to KIAA0273 [H. 31.40	
131066 AW169287 Hs.22588 ESTs 29.60 131082 Al091121 Hs.246218 ESTs; Weakly similar to zinc finger prot 9.00	
131087 AF147709 Hs.22824 ESTs; Weakly similar to p160 myb-blnding	3.86
131161 AF033382 Hs.23735 potassium voltage-gated channel; subfami 3.14 30 131179 AA171388 Hs.184482 DKFZP586D0524 protein 3.80	
30 131179 AA171388 Hs.184482 DKFZP586D0524 protein 3.80 131182 Al824144 Hs.23912 ESTs	3.67
131205 NM_003102 Hs.2420 superoxide dismutase 3; extracellular 2.98	
131277 AA131466 Hs.23767 ESTs 3.15 131281 AA251716 Hs.25227 ESTs 32.20	
35 131282 X03350 Hs.4 alcohol dehydrogenase 3 (class I); gamma	3.44
131285 Al567943 Hs.25274 ESTs; Moderately similar to putative sev 6.40 131355 R52804 Hs.25956 DKFZP564D206 protein 8.00	
131355 R52804 Hs.25956 DKFZP564D206 protein 8.00 131391 AW085781 Hs.26270 ESTs 10.00	
131461 AA992841 Hs.27263 butyrate response factor 2 (EGF-response 28.80	.03
40 131487 F13036 Hs.27373 Homo saplens mRNA; cDNA DKFZp56401763 (f 4 131517 AB037789 Hs.263395 ESTs; Highly similar to semaphorin Via [39.00	.03
131545 AL137432 Hs.28564 ESTs 11.00	
131583 AK000383 Hs.323092 ESTs; Weakly similar to dual specificity 10.00 131647 AA359615 Hs.30089 ESTs 2.47	
45 131675 H15205 Hs.30509 ESTS 3.06	
131676 Al126821 Hs.30514 ESTs 45.80 131708 S60415 Hs.30941 calcium channel: voltage-dependent: beta 2.28	
131717 X94630 Hs.3107 CD97 antigen	3.78
131756 AA443966 Hs.31595 ESTs 40.60	3.67
50 131762 AA744902 Hs.107767 ESTs; Moderately similar to CaM-KII Inhi 131821 AA017247 Hs.164577 ESTs 2.87	3.07
131839 AB014533 Hs.33010 KIAA0633 protein 3	.48
131861 AL095858 Hs.184245 KIAA0929 protein Msx2 interacting nuclea 54.00 132015 Al418006 Hs.3731 ESTs 49.20	
55 132070 BE622641 Hs.38489 ESTS 34.80	
132242 AA332697 Hs.42721 ESTs 2.68 132334 AW080704 Hs.45033 lacrimal proline rich protein 4.66	
132476 AL 119844 Hs. 49476 Homo saplens clone TUA8 Cri-du-chat regi 34.20	
132490 NM 001290 Hs 4980 NM blodfing domain 2 2.66	
60 132533 AI922988 Hs.172510 ESTs 13.00 132598 X80031 Hs.530 collagen; type IV; alpha 3 (Goodpasture 30.60	
132619 H28855 Hs.53447 ESTs; Moderately similar to kinesin ligh 4.02	
132652 N41739 Hs.61260 ESTs 3.18 132726 N52298 Hs.55608 ESTs; Weakly similar to cDNA EST yk484g1 11.43	
65 133028 R51604 Hs.300842 ESTS 2.37	
133071 BE384932 Hs.64313 ESTs 2.27 133120 NM 003278 Hs.65424 tetranectin (plasminopen-binding protein 2.63	•
133120 NM_003278 Hs.65424 tetranectin (plasminogen-binding protein 2.63 133129 AA428580 Hs.65551 ESTs	5.49
133147 AA026533 Hs.66 Interleukin 1 receptor-like 1 6.20	•
70 133151 NM_014051 Hs.94896 ESTs 3.69 · 133213 AA903424 Hs.6786 ESTs 31.40	
133276 AW978439 Hs.69504 ESTs 9.00	
133377 AJ131245 Hs.7239 SEC24 (S. cerevisiae) related gene famil 41.20 133407 AF017987 Hs.7306 secreted frizzled-related protein 1 50.20	
75 133535 AL134030 Hs.284180 protocadherin 2 (cadherin-like 2) 3.72	
133537 U41518 Hs.74602 aquaporin 1 (channel-forming integral pr	3.35
133689 NM_001872 Hs.75572 carboxypeptklase B2 (plasma) 90.80	
133779 T58486 Hs 222566 FSTs 3.05	
80 133978 AF035718 Hs.78951 transcription factor 21 2.92 133985 L34657 Hs.78146 platelet/endothelial cell adhesion molec	3.45
134000 AW175787 Hs.334841 selenium binding protein 1	4.05
134111 Al37258B Hs.8022 TU3A protein 4.49	.27
85 134204 Al873257 Hs. 7994 ESTs; Weakly shmilar to CGI-69 protein [40.80	·
t t	

	W	O 02/086	443						PCT/US	02/12476
	134641 134677	Al092634 AA251363	Hs.156114 Hs.177711	protein tyrosine phosphatase; non-recept ESTs				32.20	3.76	
	134745	NM_000685		angiotensin receptor 1B		15.00				
-	134749	T28499	Hs.89485	carbonic anhydrase IV			3.05			
5	134786	T29618	Hs.89640	angiopoietin 1 receptor; TEK tyrosine ki				57.80		3.73
	134825	U33749	Hs.197764	thyroid transcription factor 1			0.50			3.73
	134978 135010	A1829008 N50465	Hs.333383 Hs.92927	ficelin (collagen/fibrinogen domain-cont ESTs			2.52	31.60		•
	135053	AW796190	Hs.93678	ESTs				31.00	3.21	
10	135081	AF069517	Hs.173993	RNA binding motif protein 6	28.80				0.2.	
	135091	AA493650	Hs.94367	ESTs						4.24
	135135	AA775910	Hs.95011	syntrophin; beta 1 (dystrophin-associate		8.00				
	135203	C15737	Hs.269386	ESTs					4.31	
16	135236	AI636208	Hs.96901	ESTs	43.00				_	**
15	135266	R41179	Hs.97393	Human mRNA for KIAA0328 gene; partial cd					6.	.42
	135346	NM_000928	Hs.992	phospholipase A2; group iB (pancreas)			3.82			
	135378	AW961818	Hs.24379	potassium voltage-gated channel; shaker-	37.20		4.15			
	135387 135388	NM_001972 W27965	Hs.99863 Hs.99865	elastase 2; neutrophil EST	38.80					
20	135402	L12398	Hs.99922	dopamine receptor D4	30.00				4.21	

TABLE 2B shows the accession numbers for those primekeys lacking unigenelD's for Table 2A. For each probeset we have listed the gene cluster number from which the oligonucleotides were designed. Gene clusters were compiled using sequences derived from Genbank ESTs and mRNAs. These sequences were clustered based on sequence similarity using Clustering and Alignment Tools (DoubleTwist, Oakland California). The Genbank accession numbers for sequences comprising each cluster are listed in the "Accession" column.

	Pkev:	Unique Eos probeset identif	ier number
	CAT numb	per. Gene cluster number	
30	Accession	: Genbank accession number	3
	Pkey	CAT number Accessions	
35	108447 108550	434527 AA079126 120073_1 AA084867	A A DE A POR
33	108655	120073_1 AA0099960 /	
	102397	443711 U41898	WI 13013
	126303	1525933_1 D78841 D7	oogn
	125810	1554054_1 H00083 R8	
40	103627	2615_2 Z48513 Z48	
40	121366		A405617 AW276706
	114609	116777_1 AA079505	
	115272		AA211890 AA279425
	108338	112186_1 AA070773	
45	108434		AQ078782 AAQ75788
	123802		AA620448
	102310	NOT FOUND entrez U3383	
	102636	entrez U67092 U67092	
	104776	genbank_AA026349	AA026349
50	120504		AA256837
	113502	genbank_T89130T89130	
	108499		AA083103
	101308	entrez_L41390 L41390	
	108629		AA102425
55	103098	221_215 M86361 Z26	5593 X02850 D13070 AE000559 M17649 M87869 M87871 X61077 M16286 AF018169 X61079 S59351 X60142 AF043169
	103241	entrez_X76223 X76223	
	103508	entrez_Y10141 Y10141	
	103575	entrez_Z26256 Z26256	
~	119514	NOT_FOUND_entrez_W3793	
60	121082		AA398722
	128634	AA464918_at AA464918	
	105817		AA397825
	121518		AA412155
CE	114449		AA020736
65	114648		AA101056
	121950		AA429515
	107723	genbank_AA015967	AA015967

Table 3A shows 452 genes up-regulated in chronically diseased lung relative to normal lung. Chronically diseased lung samples represent chronic non-malignant lung diseases such as fibrosis, emphysema, and bronchitis. These genes were selected from 59680 probesels on the Eos/Affymetrix Hu03 Genechip array. Gene expression data for each probesel obtained from this analysis was expressed as average intensity (AI), a normalized value reflecting the relative level of mRNA expression.

5	Pkey:	Unique Eos probeset identifier number
	ExAccn:	Exemplar Accession number, Genbank accession number
	UnigenelD:	Unigene number
	Unigene Tille:	Unigene gene title
	R1:	80th percentile of AI for chronically diseased lung samples divided by the 90th percentile of AI for normal lung samples.
10	R2:	80th percentile of AI for chronically diseased lung samples divided by the 90th percentile of normal lung samples, squamous cell carcinomas and
		adenocarcinomas
	R3:	70th percentile of AI for chronically diseased lung samples minus the 15th percentile of AI for all normal lung, chronically diseased lung and tumor samples
		divided by the 90th percentile of normal lung samples, squamous cell carcinomas and adenocarcinomas minus the 15th percentile of Al for all normal lung,
		chronically diseased lung and tumor samples
15		

Pkey	ExAccn	UnigenelD	Unigene Title	R1	R2	R3
135423	U50531	Hs.138751	Human BRCA2 region, mRNA sequence CG030	12.40		2.13
135346	NM_000928	Hs.992	phospholipase A2, group IB (pancreas)	12.40		
135057	U90268	Hs.93810	cerebral cavernous malformations 1	11.67		
134951 134799	BE305081 M36821	Hs.169358 Hs.89690	hypothetical protein GRO3 oncogene		8.00 8.20	
134786	T29618 NM 000829	Hs.89640 Hs 163697	TEK tyrosine kinase, endotheliai (venous plutamate receptor, ignotrophic, AMPA 4	29.80		
134752	BE246762	Hs.89499	arachidonate 5-lipoxygenase			1.93 2.07
134698	BE326276	Hs.8861	ESTs	40.00		2.07
134636 134627	NM_005582 Al018768	Hs.12482	glyceronephosphate O-acyltransferase	13.60		1.92
134622 134570	AW975159 U66615			13.20		1.92
134561	U76421	Hs.85302	adenosine deaminase, RNA-specific, B1 (h		6.20	1.78
134417	NM_006416	Hs.82921	solute carrier family 35 (CMP-siatic aci		O,LO	
134343 134323	D50683 BE170651	Hs.82028 Hs.8700	deleted in liver cancer 1			
134300 134299	NM_001430 AW580939	Hs.8136 Hs.97199	endothellal PAS domain protein 1 complement component C1g receptor			
134253	X52075	Hs.80738	sialophorin (gpL115, leukosialin, CD43)	20.60 12.20		
133985	L34657	Hs.78146	platelet/endothelial cell adhesion molec	,		
133835	A1677897	Hs.76640	RGC32 protein			
133651 133633	Al301740 D21262	Hs.173381 Hs.75337	dihydropyrimidinase-like 2 nucleotar and colled-body phosphprotein	15.20		
133565 133548	AW955776 AW946384	Hs.313500 Hs.178112	ESTs, Moderately similar to ALU7_HUMAN A DNA segment, single copy probe LNS-CAI/L			1.77
133488	AA335295	Hs.74120	adipose specific 2			2.08
133337	AF085983	Hs.293676	ESTs		9.60	1.77
133153	AF070592	Hs.66170	HSKM-B protein	30.60		1.77
133130 133120	AI128606 NM_003278	Hs.6557 Hs.65424	zinc finger protein 161 tetranectin (plasminogen-binding protein			
132928 132836	AW168082 AR023177	Hs.169449 Hs.29900	protein kinase C, alpha KIAA0960 protein	13.80		
132799	W73311	Hs.169407	SAC2 (suppressor of actin mutations 2,	41.60 40.40		
132548	X12830	Hs.193400	interleukin 6 receptor	70,70	7.20	
132439	AK001942	Hs.4863	hypothetical protein DKFZp566A1524		4.70	1.88
132240 132210	AB018324 NM_007203	Hs.42676 Hs.42322	KIAA0781 protein A kinase (PRKA) anchor protein 2	21.20		1.99
132199	AL041299 T96555	Hs.165084 Hs.31562	ESTs ESTs	15.20		1.76
131745	Al828559	Hs.31447	ESTs, Moderately similar to A46010 X-li	27.80	4.00	
131686	NM_012296	Hs.30687	GRB2-associated binding protein 2		•	
131676 131629	A1126821 Z45794	Hs.238809	ESTs	21.40	0.20	
131589 131536	C18825 AA019201		ESTs		9.40	
131517	AB037789	Hs.263395	sema domain, transmembrane domain (TM), DKFZP564D206 protein		3.59 4.48	
131253	R71802	Hs.24853	ESTs	15.00		1.75
131156	A1472209	Hs.323117	ESTs		254	1.84
131066 131061	N64328	Hs.268744	KIAA1796 protein		3.54	
131053 130895	AA348541 AA641767	Hs.296261 Hs.21015	guanine nucleotide binding protein (G pr hypothetical protein DKFZp564L0864 simil	16.60		1.93
130762	D84371	Hs.1898	paraoxonase 1	12.00		
	135423 135378 135378 13536 135235 135235 135235 134799 134752 134752 134752 134636 134627 134521 134531 134323 134323 134323 134329 134253 134329 134263 133985 133985 133985 133985 133985 133985 133985 133985 133985 133985 133985 133985 133985 133558 133558 133558 133558 133558 133579 132240 132210 132298 132240 132210 132298 132199 131751 131676 131626 131636	135423 U50531 135378 AW9561818 135378 AW9561818 135346 NM_000928 135235 W296244 135057 U90268 134799 M36821 134786 T29618 134772 NM_000829 134752 BE246762 134749 T28499 134636 NM_005582 134637 Al018768 134622 AW975159 134561 U76421 134468 NM_001772 134417 NM_006416 134343 D50683 134523 BE170651 134501 WM_001430 134299 AW580939 134253 X52075 134182 D52059 133978 AF035718 133835 Al677897 133978 AF035718 133835 Al677897 133651 Al301740 133633 D21262 133565 AW955776 133548 AW946384 133488 AA335295 133478 AW946384 133489 AW946384 133489 AW946384 133489 AW946384 133480 AW9465880 133150 AW960246 131660 AW169287 131656 AW71209 131551 R52804 131553 R52804 131553 R71802 131656 AW71209 131656 AW71209 131656 AW716206 131666 AW716208 131666 AW716208 131666 AW716209 131667 AM3684541 1306995 AA641767	135423 U50531 Hs.138751 135378 AW961818 Hs.24379 135346 NM_000928 Hs.992 135235 AW298244 Hs.293507 135057 U90268 Hs.99310 134795 BE305081 Hs.163587 134799 M36821 Hs.89640 134772 NM_000829 Hs.163597 134752 BE246762 Hs.89499 134752 BE246762 Hs.89499 134762 AW975159 Hs.946361 134636 NM_005582 Hs.293097 134570 U66615 Hs.87205 134627 AU975159 Hs.12482 134627 AU975159 Hs.12482 134628 NM_001772 Hs.83731 134570 NM_0008416 Hs.83032 134461 NM_0016416 Hs.83032 134461 NM_0016416 Hs.83031 134417 NM_006416 Hs.83031 134340 NM_001430 Hs.8136 134299 AW580939 Hs.7972 134530 NM_001430 Hs.8136 133835 AI677897 Hs.780640 133851 AI677897 Hs.76640 133651 AI301740 Hs.78061 133835 AI677897 Hs.76640 133651 AI301740 Hs.76640 133651 AI301740 Hs.76640 133651 AI301740 Hs.76640 133631 AI2666 Hs.76640 133634 AW945348 Hs.78061 133130 AI126606 Hs.76640 133130 AI126606 Hs.65570 133135 AF070592 Hs.66170 132742 AA025480 Hs.293676 132299 AU041299 Hs.66570 131554 AV940324 Hs.293676 131556 AW001942 Hs.4663 132240 AB018324 Hs.29506 131557 AB037789 Hs.31650 131558 AV001942 Hs.26321 131556 AV001942 Hs.26326 131253 R71802 Hs.26326 131556 AV601929 Hs.263367 131556 AV601929 Hs.263367 131556 AV601929 Hs.263367 131556 Hs.26326 131556 Hs.86363 132210 NM_0012296 Hs.263367 131556 Hs.86363 132210 NM_0012296 Hs.263363 131551 Hs.26321 131551 Hs.26321 131551 Hs.26321 131551 Hs.26321 131552 Hs.26326 131553 Hs.26326 131553 Hs.26326 131553 Hs.26326 131553 Hs.26326 131553 Hs.26326 131554 Hs.26326 131555 Hs.36363	135423 U50531 Hs. 138751 135378 AW561818 Hs. 24379 135348 NM_000928 Hs. 2992 136253 NW298244 Hs. 29557 136257 U90268 Hs. 29350 13799 M36821 Hs. 89890 13479 M36821 Hs. 89890 134772 NM_000829 Hs. 163597 134752 BE246762 Hs. 89499 134636 NM_005582 Hs. 88705 134636 NM_005582 Hs. 88705 134623 AW576159 Hs. 87205 134624 NW576159 Hs. 87205 134651 U76421 Hs. 85302 134561 U76421 Hs. 85302 134468 NM_001772 Hs. 83731 134461 NM_008681 Hs. 829281 134330 NM_001430 Hs. 822921 134330 NM_001430 Hs. 82293 134329 AW580393 Hs. 77932 134622 AW7869383 Hs. 77897 133997 AF085983 Hs. 77972 133986 AG37789 Hs. 76383 13320 AB037715 Hs. 78180 133337 AF085983 Hs. 774120 133337 AF085983 Hs. 774120 133337 AF085983 Hs. 78120 133337 AF085983 Hs. 78132 133330 AL18660 Hs. 183639 133320 AB037715 Hs. 86557 133120 NM_003248 Hs. 85572 133774 AW005484 Hs. 29850 131553 AR019201 Hs. 829317 131563 AR019201 Hs. 82951 131563 AR019201 Hs. 829317 131563 AR019201 Hs. 829317 131563 AR019201 Hs. 829317 131563 AR019201 Hs. 829317 131563 AR019201 Hs. 829357 131563 AR	135423	184623

	W	O 02/086	443				
	130657	AW337575	Hs.201591	ESTs			
	130655	AI831962	Hs.17409	cysteine-rich protein 1 (intestinal)			2.08
	130589 130562	AL110226 D50402	Hs.16441 Hs.182611	DKFZP434H204 protein solute carrier family 11 (proton-coupled			1.91
5	130555	R69743	Hs.116774	Integrin, alpha 1		9.60	
-	130365	W56119	Hs.155103	eukaryotic translation initiation factor	11.60		
	130273	AW972422	Hs.153863	MAD (mothers against decapentaplegic, Dr		6.60	4.04
	130259	NM_000328	Hs.153614 Hs.132390	retinitis pigmentosa GTPase regulator	21.20		1.91
10	130090 129958	H97878 R27496	Hs.1378	zinc finger protein 36 (KOX 18) annexin A3	21.20	5.05	
	129898	AI672731	Hs.13256	ESTs		•	
	129875	AA181018	Hs.13056	hypothetical protein FLJ13920	18.60		
	129699	AB007899	Hs.12017	homolog of yeast ubiquitin-protein ligas			
15	129626 129598	F13272 N30436	Hs.111334 Hs.11556	ferritin, light polypeptide Homo sapiens cDNA FLJ12566 fis, clone NT	22.63		
13	129593	AI338247	Hs.98314	Homo sapiens mRNA; cDNA DKFZp586L0120 (f	22.00		
	129565	X77777	Hs.198726	vasoactive intestinal peptide receptor 1			2.53
	129527	AA769221	Hs.270847	delta-tubulin	39.20		0.44
20	129402	W72062	Hs.11112	ESTs	15.20		2.11
20	129385 129315	AA172106 NM_014563	Hs.110950 Hs.174038	Rag C protein spondyloepiphyseal dysplasia, late	12.40		
	129312	T97579	Hs.110334	ESTs, Weakly similar to 178885 serine/th	20.83		
	129240	AA361258	Hs.237868	interleukin 7 receptor			1.95
25	129210	AL039940	Hs.202949	KIAA1102 protein		4.20	
23	129122 129057	AW958473 N90866	Hs.301957 Hs.276770	nudix (nucleoside diphosphate linked moi CDW52 antigen (CAMPATH-1 antigen)		4.20	
	128946	Y13153	Hs.107318	kynurenine 3-monooxygenase (kynurenine 3		5.20	
	128798	AF015525	Hs.302043	chemokine (C-C motif) receptor-like 2			
20	128789	AW368576	Hs.139851	caveolin 2	40.00		2.24
30	128778 128766	AA504776 AW160432	Hs.186709 Hs.296460	ESTs, Weakly similar to 138022 hypothet craniofacial development protein 1	12.20 26.40		
	128631	R44238	Hs.155546	KIAA1080 protein; Golgi-associated, gamm	20.70		1.78
	128624	BE154765	Hs.102647	ESTs, Weakly similar to TRHY_HUMAN TRICH			2.51
25	128609	NM_003616	Hs.102456	survival of motor neuron protein interac	16.00		
35	128603	NM_004915	Hs.10237	ATP-binding cassette, sub-family G (WHIT potassium inwardly-rectifying channel, s	12.80	4.00	
	128598 128458	AA305407 H55864	Hs.102308 Hs.56340	ESTs		4.00	
	128061	AF150882	Hs.186877	sodium channel, voltage-gated, type XII,	17.20		
40	127968	AA830201	Hs.124347	ESTs	21.30		
40	127959	Al302471	Hs.124292	Homo sapiens cDNA: FLJ23123 fis, clone L	40.60		
	127944 127925	Al557081 AA805151	Hs.262476 Hs.3628	S-adenosylmethlonine decarboxylase 1 mitogen-activated protein kinase kinase	10.60 13.40		
	127896	A1669586	Hs.222194	ESTs		7.00	
45	127859	AA761802	Hs.291559	ESTs	14.00		
45	127817	AA836641	Hs.163085	ESTs	14.00		
	127742 127628	AW293496 Al240102	Hs.180138 Hs.322430	ESTs NDRG family, member 4	11.00 11.10		
	127609	X80031	Hs.530	collagen, type IV, alpha 3 (Goodpasture			
50	127582	AA908954	Hs.130844	ESTs	19.60		
50	127543	AK000787	Hs.157392	Homo saplens cDNA FLJ20780 fis, clone CO	15.40 17.50		
	127535 127404	AA568424 AJ379920	Hs.164450 Hs.270224	ESTs ESTs	14.60		
	127396	L31968	Hs.187991	DKFZP564A122 protein	15.40		
<i></i>	127374	AA442797	Hs.312110	ESTs, Weakly similar to 138022 hypothet	14.60		
55	127346	AA203616	Hs.44896	DnaJ (Hsp40) homolog, subfamily 8, membe	21.00		
	127340 127307	BE047653 AW962712	Hs.119183 Hs.126712	ESTs, Weakly similar to ZN91_HUMAN ZINC ESTs, Weakly similar to AF191020 1 E2IG5	15.80		
	127242	AW390395	Hs.181301	cathepsin S	22.60		
C O	127167	AA625690	Hs.190272	ESTs	21.40		
60	127046	AA321948	Hs.293968	ESTs	41.20 11.00		
	126928 126900	AA480902 AF137386	Hs.137401 Hs.12701	EST8 plasmolipin	11.00		1.78
	126852	AA399961		gb:zu68c01.r1 Soares_testis_NHT Homo sap		5.60	
<i>C</i> =	126816	AA248234		gb:csg2228.seq.F Human fetal heart, Lamb	12.20		
65	126812	AB037860	Hs.173933	nuclear factor I/A	17.19		
	126666 126645	AA648886 AA316181	Hs.151999 Hs.81635	ESTs six transmembrane epithelial antigen of	13.57 15.40		
	126592	AI611153	Hs.6093	Homo saplens cDNA: FLJ22783 fis, clone K	101.10	4.67	
70	126556	AF255303	Hs.112227	membrane-associated nucleic acid binding	18.00		
70	126433	AA325606	U- 000075	gb:EST28707 Cerebellum II Homo sapiens c	16.77		
	126299 126218	AW979155 AL049801	Hs.298275 Hs.13649	amino acid transporter 2 Novel human gene mapping to chomosome 13	14.60	3.50	
	126182	AA721331	Hs.293771	ESTs	13.40	0.00	
~~	126177	AW752782	Hs.129750	hypothetical protein FLJ10546	18.20		
75	126142	H86261	Hs.40568	ESTs	14.00		
	126077 125994	M78772 Al990529	Hs.210836 Hs.270799	ESTs ESTs	16.59 17.40		
	125934	AA193325	Hs.32646	hypothetical protein FLJ21901	13.00		
00	125847	AW161885	Hs.249034	ESTs	49.57		
80	125831	H04043	11- 000-0	gb:yj45c03.r1 Soares placenta Nb2HP Homo	42.00		
	125731	R61771 BE612918	Hs.26912 He 151073	ESTs hypothetical protein FLJ23511	13.20 11.20		
	125676 125561	F18572	Hs.151973 Hs.22978	ESTs, Weakly similar to ALU4_HUMAN ALU S	1120		
0.5	125552	H09701	Hs.278366	ESTs, Weakly similar to 138022 hypotheti	12.60		
85	125489	H49193	Hs.124984	ESTs, Moderately similar to ALU7_HUMAN A	33.40		

	W	O 02/086	443				
	125422	AA903229	Hs.153717	ESTs	20.00		1.80
	125331	Al422996	Hs.161378	ESTS	38.00 18.20		
	125309 125167	T12411 AL137540	Hs.183745 Hs.102541	hypothetical protein FLJ13456 netrin 4	10.20		1.95
5	125139	AW194933	Hs.9788	hypothetical protein MGC10924 similar to			1.84
_	125042	T78906	Hs.269432	ESTs, Moderately similar to ALU1_HUMAN	21.80		
	124711	NM_004657	Hs.26530	serum deprivation response (phosphatidy)	23.20	10.60	
	124631 124578	NM_014053 N68321	Hs.270594 Hs.231500	FLVCR protein EST	21.43		
10	124574	AL036596	Hs.42322	A kinase (PRKA) anchor protein 2			1.77
	124472	N52517	Hs.102670	EST	37.20		
	124438	BE178536	Hs.11090	membrane-spanning 4-domains, subfamily A	14.64		
	124357 124308	N22401 AW973078	Hs.293039	gb:yw37g07.s1 Morton Fetal Cochlea Homo ESTs	14.04	4.00	
15	124300	H58608	Hs.151323	ESTs			
	124097	AW298235	Hs.101689	ESTs		27.20	
	123978	T89832	Hs.170278	ESTs		6.00	2.03
	123972 123961	T46848 AL050184	Hs.70337 Hs.21610	Immunoglobulin superfamily, member 4 DKFZP434B203 protein		0.00	1.79
20	123936	NM_004673	Hs.241519	angiopoletin-like 1		15.80	
	123802	AA620448	***************************************	gb:ae58c09.s1 Stratagene lung carcinoma		4.23	
	123734	AA609861	Hs.312447	ESTs	22.00	4.20	
	123619	AA602964	Hs.112640	gb:no97c02.s1 NCI_CGAP_Pr2 Homo sapiens EST	33.60 10.93		
25	123596 123476	AA421130 AA384564	Hs.108829	ESTs			2.18
	123340	AA504264	Hs.182937	peptidylprolyl isomerase A (cyclophilin	11.20		
	123190	AA489212	Hs.105228	EST	14.20	7.00	
	123136 123073	AW451999 AA485061	Hs.194024 Hs.105652	ESTs ESTs	31.20	7.00	
30	123055	AA482005	Hs.105102	ESTs, Weakly similar to reverse transcri		4.80	
	122699	AA456130	Hs.301721	KIAA1255 protein	44.40	5.00	
	122679	AA811286	Hs.192837	ESTs, Wealdy similar to ALU5_HUMAN ALU S	14.40		
	122633 122553	NM_001546 AA451884	Hs.34853 Hs.190121	inhibitor of DNA binding 4, dominant neg ESTs	40.00		
35	122544	AW973253	Hs.292689	ESTs	15.40		
	122485	AA524547	Hs.160318	FXYD domain-containing ion transport reg		40.40	1.81
	122211	AA300900	Hs.98849	ESTs, Moderately similar to AF161511 1 H		12.10	1.95
	122127 122011	AW207175 AA431082	Hs.106771	ESTs gb:zw78a10.s1 Soares_testis_NHT Homo sap			1.89
40	121992	AI860775	Hs.98506	ESTs		3.60	
	121989	W56487	Hs.193784	Homo sapiens mRNA; cDNA DKFZp586K1922 (f			2.01 1.85
	121835 121726	AB033030 AF241254	Hs.300670 Hs.178098	KIAA1204 protein angiotensin I converting enzyme (peptidy	12.43		1.00
	121690	AV660305	Hs.110286	ESTs			1.82
45	121643	AA640987	Hs.193767	ESTs	14.00		
	121633 121622	AA417011 AA416931	Hs.98175 Hs.126065	EST ESTs	14.00	16.40	
	121497	AA412031	Hs.97901	EST	11.20		
50	121351	AW206227	Hs.287727	hypothetical protein FLJ23132	12.20		1.83
50	121314 121242	W07343 AA400857	Hs.182538 Hs.97509	phospholipid scramblase 4 ESTs	22.40		1.03
	121059	AA400657 AA393283	па.э/ 509	gb:zt74e03.r1 Soares_testis_NHT Homo sap	14.80		
	120934	AA226198		gb:nc26a07.s1 NCI_CGAP_Pr1 Homo sapiens	21.20		
<i>5</i>	120755	AA312934	Hs.190745	Homo saplens cDNA: FLJ21326 fis, clone	20.00		1.79
55	120637 120484	AA811804 AA253170	Hs.96473	gb:ob39a05.s1 NCI_CGAP_GCB1 Homo saplens EST	40.20		
	120336	N85785	Hs.181165	eukaryotic translation elongation factor	75	6.60	
	120266	AI807264	Hs.205442	ESTs, Weakly similar to T34036 hypotheti	16.80	. 70	
60	120132	W57554	Hs.125019	ESTs ESTs		4.73	1.75
00	120041 119996	AA830882 W88996	Hs.59368 Hs.59134	EST		7.20	.,,,
	119970	AA767718	Hs.93581	hypothetical protein FLJ10512	11.20		
	119861	W78816	Hs.49943	ESTs, Weakly similar to S65657 alpha-1C-		3.78	
65	119824 119740	W74536 AW021407	Hs.184 Hs.21068	advanced glycosylation end product-spect hypothetical protein	20.20		
05	119271	A1061118	Hs.65328	Fanconi anemia, complementation group F	15.20		
	119221	C14322	Hs.250700	tryptase beta 1	40.00		
	119126	R45175	Hs.117183	ESTs	12.60		
70	119073 118928	BE245360 AA312799	Hs.279477 Hs.283689	ESTs activator of CREM in testis		10.00	
, 0	118901	AW292577	Hs.94445	ESTs		3.96	
	118661	AL137554	Hs.49927	protein kinase NYD-SP15	10.10	9.60	
	118607	Al377444 ·	Hs.54245	ESTs, Weakly similar to S65824 reverse thypothetical protein FLJ21939 similar to	10.40		1.90
75	118449 118416	AI813865 N66028	Hs.164478 Hs.49105	FKBP-associated protein	16.20		
	118379	N64491	Hs.48990	ESTs		4.00	
	118329	N63520		gb:yy62f01.s1 Soares_multiple_sclerosis_		6.60	
	118320 118253	N63451 AA497044	Hs.141600 Hs.20887	ESTs, Weakly similar to alternatively s hypothetical protein FLJ10392	17.60	3.80	
80	118124	N56968	Hs.46707	chromosome 21 open reading frame 37	14.00		
	118056	AB037746	Hs.42768	hypothetical protein DKFZp76100113		F 44	1.86
	118032	N52802	Hs.47544	EST Homo sapiens clone 23632 mRNA sequence		5.00 4.00	
	117840 117404	T26379 N39725	Hs.48802 Hs.15220	zinc finger protein 106		7.00	1.90
85	117314	N32498	Hs.42829	ESTs	14.20		

	W	O 02/086	443				
	117209	W03011	Hs.306881	MSTP043 protein			
	117023	AW070211	Hs.102415	Homo sapiens mRNA; cDNA DKFZp586N0121 (f			2.31
	116814	H50834		gb:yp86a10.s1 Soares fetal fiver spleen	20.20		
_	116784	AB007979	Hs.301281	Homo sapiens mRNA, chromosome 1 specific		3.51	
5	116766	A1608657	Hs.95097	ESTs	16.20		
	116712	AW901618	Hs.61935	Homo saplens mRNA; cDNA DKFZp761l071 (fr		6.80	
	116707	H10344	Hs.49050	ESTs, Weakly similar to A Chain A, Human	18.60		
	116351	AL133623	Hs.82501	similar to mouse Xm1 / Dhm2 protein	19.40		
10	116279	AW971248	Hs.291289	ESTs, Weakly similar to ALU1_HUMAN ALU S			2.42
10	116166	AL039940	Hs.202949	KIAA1102 protein			2.13 1.75
	116152	AL040521	Hs.15220	zinc finger protein 106	13.20		1.75
	116117 116107	BE613410	Hs.31575	SEC63, endoplasmic reticulum translocon hypothetical protein FLJ20093	30.11		
		AL133916 AA001732	Hs.172572 Hs.173233	hypothetical protein FLJ10970	30.11		2.36
15	115965 115955		Hs.44198	intracellular membrane-associated calciu	18.20		2.00
13	115844	AF263613 Al373062	Hs.332938	hypothetical protein MGC5370	18.57		
	115683	AF255910	Hs.54650	iunctional adhesion molecule 2	10.57	23.00	
•	115673	AA406341	Hs.269908	Homo sapiens cDNA FLJ11991 fls, clone HE	11.82	20.00	
	115672	Al889110	Hs.73251	ESTs	10.60		
20	115566	AJ142336	Hs.43977	Human DNA sequence from clone RP11-196N1	10.00		1.76
	115313	AA808001	Hs.184411	albumin	25.20		
	115279	AW964897	Hs.290825	ESTs		8.00	
	115230	AA278300	Hs.124292	Homo saplens cDNA: FLJ23123 fis, done L			1.80
	115110	AK001671	Hs.11387	KIAA1453 protein	14.20		
25	114999	BE246481	Hs.87856	ESTs	19.20		
	114930	AA237022	Hs.188717	ESTs		5.60	
	114922	AA235672	Hs.87491	ESTs		3.60	
	114837	BE244930	Hs.166895	ESTs	43.70		
20	114769	AA149060	Hs.296100	ESTs	11.00		
30	114761	AA143781	Hs.126280	hypothetical protein FLJ23393	14.00		
	114736	AI610347	Hs.103812	ESTs, Moderately similar to ALU1_HUMAN A		4.20	
	114596	AA310162	Hs.169248	cytochrome c	10.71		
	114518	AW163267	Hs.106469	suppressor of var1 (S.cerevisiae) 3-like	20.40		
35	114455	H37908	Hs.271616	ESTs, Weakly similar to ALUS_HUMAN ALU S	20.40	47.00	
22	114452	Al369275	Hs.243010	Homo saplens cDNA FLJ14445 fis, clone HE		17.20	2.00
	114359	NM_016929	Hs.283021	chloride intracellular channel 5	12.40		2.09
	114357 114251	R41677	Hs.6107	Homo sapiens cDNA FLJ14839 fis, clone OV ESTs	12.40		2.00
	114138	H15261 AW384793	Hs.21948 Hs.15740	Homo sapiens mRNA; cDNA DKFZp434E033 (fr		11.40	2.00
40	114124	W57554	Hs.125019	ESTS		6.04	
70	113946	AW083883	Hs.37896	Homo sapiens cDNA FLJ13510 fis, clone PL		0.04	1.82
	113695	T96965	Hs.17948	ESTs, Weakly similar to ALUB_HUMAN IIII			1,02
	113606	NM_013343	Hs.278951	NAG-7 protein			2.15
	113590	R49642	Hs.142447	ESTs, Weakly similar to ALU1_HUMAN ALU S		3.60	
45	113560	T91015	Hs.268626	ESTs	32.00	••••	
	113552	AI654223	Hs.16026	hypothetical protein FLJ23191			
	113540	AW152618	Hs.16757	ESTs			
	113502	T89130		gb:ye12d01.s1 Stratagene lung (937210) H		8.35	
	113288	A1076838	Hs.12967	ESTs ·	12.40		
50	113252	NM_004469	Hs.11392	c-fos induced growth factor (vascular en		4.27	
	113238	R45467	Hs.189813	ESTs			
	113203	AA743563	Hs.10305	ESTs	21.20		
	113195	H83265	Hs.8881	ESTs, Weakly similar to S41044 chromosom	44.00		1.92
55	113089	T40707	Hs.270862	ESTs	14.33	0.00	
55	113076	AF033199	Hs.8198	zinc finger protein 204		6.00	
	113009	T23699	Hs.7246 Hs.6295	ESTs		9.40 12.20	
	112937 112891	A1694320 T03927	Hs.293147	ESTs, Weakly similar to T17248 hypotheti ESTs, Moderately similar to A46010 X-li	10.57	12.20	
	112794	R97018	110.230177	gb:yq74b08.s1 Soares fetal liver spleen	28.60		
60	112691	R88708	Hs.220647	ESTs	15.33		
	112602	AW004045	Hs.203365	ESTs	15.60		
	112366	AF035318	Hs.12533	Homo saplens clone 23705 mRNA sequence	15.40		
	112210	R49645	Hs.7004	ESTs	14.00		
	112064	AL049390	Hs.22689	Homo saplens mRNA; cDNA DKFZp586O1318 (f	13.00		
65	111998	R42379	Hs.138283	ESTs	11.00		
	111987	NM_015310	Hs.6763	KIAA0942 protein	22.40		
	111803	AA593731	Hs.325823	ESTs, Moderately similar to ALU5_HUMAN A			1.77
	111737	H04607	Hs.9218	ESTs			1.86
70	111605	T91061	Hs.194178	ESTs, Moderately similar to PC4259 ferri	23.00		
70	111510	R07856	Hs.16355	ESTs	11.02		
	111341	AL157484	Hs.22483	Homo sapiens mRNA; cDNA DKFZp762M127 (fr	40 40		1.88
	111280	AA373527	Hs.19385	CGI-58 protein	18.40		
	111247	AW058350	Hs.16762	Homo sapiens mRNA; cDNA DKFZp564B2062 (f	27 50		
75	111232	AI247763	Hs.16928	ESTs ESTs	27.60 14.80		
, ,	110942 110924	R63503 AW058463	Hs.28419 Hs.12940	zinc-fingers and homeoboxes 1	24.71		
	110837	H03109	Hs.108920	HT018 protein	47.7		2.18
	110824	Al767183	Hs.26942	ESTs	12.20		Z. 10
_	110776	AB032417	Hs.19545	frizzled (Drosophila) homolog 4			1.75
80	110576	H60869	Hs.37889	ESTs	13.00		•
	110369	AK000768	Hs.107872	hypothetical protein FLJ20761		5.60	
		R44557	Hs.23748	ESTs		-	2.31
	110099	1441001					2.0
	10099	A1796320	Hs.10299	Homo sapiens cDNA FLJ13545 fis, clone PL			2.01
0.5		A1796320 AA001266		Homo septens cDNA FLJ13545 fis, clone PL ESTs	11.25		
85	109984	A1796320	Hs.10299	Homo sapiens cDNA FLJ13545 fis, clone PL	11.25		2.68

	W	O 02/086	443				
	109842	AW818436	Hs.23590	solute carrier family 16 (monocarboxylic	23.83		
	109837	H00656	Hs.29792	ESTs, Weakly similar to 138022 hypotheti		47.00	3.91
	109796	A1800515	Hs.12024	ESTs		17.20	
5	109688	R41900	Hs.22245	ESTS	22.80	9.60	
3	109648	H17800	Hs.7154 Hs.27519	ESTs ESTs	22.00		
	109613 109550	H47315 AW021488	Hs.26981	ESTs			
	109523	AW193342	Hs.24144	ESTs			1.89
	109472	AK001989	Hs.91165	hypothetical protein		6.00	
10	109355	AA524525	Hs.48297	DKFZP586C1620 protein	15.00		
	109260	AW978515	Hs.131915	KIAA0863 protein	25.60		
	108781	AA128654		gb:zn98g07.s1 Stratagene fetal retina 93	14.20		
	108663	BE219231	Hs.292653	ESTs, Weakly similar to T26845 hypotheti	11.00 26.00		
15	108573 108480	AA086005 AL133092	Hs.68055	gb:zl84c04.s1 Stratagene colon (937204) hypothetical protein DKFZp434l0428	20.00		
13	108382	NM_006770	Hs.67726	macrophage receptor with collagenous str			1.83
	108174	AA055632	Hs.303070	ESTs	15.20		
	108138	AL049990	Hs.51515	Homo sapiens mRNA; cDNA DKFZp564G112 (fr		3.60	
	108087	AA045708	Hs.40545	ESTs	15.44		
20	108048	AI797341	Hs.165195	Homo sapiens cDNA FLJ14237 fis, clone NT		11.40	
	108041	AW204712	Hs.61957	ESTs		A 70	
	107997	AL049176	Hs.82223	chordin-like		4.76	
	107994 107922	AA036811 BE153855	Hs.48469 Hs.61460	LIM domains containing 1 Ig superfamily receptor LNIR	14.20		
25	107522	BE379594	Hs.49136	ESTs, Moderately similar to ALU7_HUMAN A	51.80		
	107666	AA010611	Hs.60418	EST	29.20		
	107332	T87750	Hs.183297	DKFZP566F2124 protein	10.73		
	107292	BE166479	Hs.4789	Homo sapiens serologically defined breas	32.00		
20	107230	A1034467	Hs.34650	ESTs	17.40		
30	107168	W57578	Hs.237955	RAB7, member RAS encogene family	10.43		
	107160	AA314490	Hs.27669	KIAA1563 protein KIAA1272 protein	11.40		
	107054 107029	AI076459 AF264750	Hs.15978 Hs.288971	myelold/lymphold or mixed-lineage leukem	21.40		
	106999	H93281	Hs.10710	hypothelical protein FLJ20417	35.80		
35	106954	AF128847	Hs.204038	indolethylamine N-methyltransferase			1.76
-	106870	Al983730	Hs.26530	serum deprivation response (phosphatidy)			
	106865	AW192535	Hs.19479	ESTs	13.40		
	106844	AA485055	Hs.158213	sperm associated antigen 6		7.13	
40	106820		· Hs.12592	period (Drosophila) homolog 3	40.00	7.00	
40	106818	AK002135	Hs.3542	hypothetical protein FLJ11273	13.00		2.05
	106797	A1768801	Hs.169943 Hs.188833	Homo sapiens cDNA FLJ13569 fis, clone PL ESTs			2.00
	106773 106747	AA478109 NM_007118	Hs.171957	triple functional domain (PTPRF interact	12.60		
	106743	BE613328	Hs.21938	hypothetical protein FLJ12492	10.60		
45	106667	AW360847	Hs.16578	ESTs			
	106605	AW772298	Hs.21103	Homo sapiens mRNA; cDNA DKFZp564B076 (fr			2.40
	106567	AW450408	Hs.86412	chromosome 9 open reading frame 5			1.78
	106562	AL031846	Hs.152151	plakophilin 4			1.76 2.19
50	106536 106533	AA329648 AL134708	Hs.23804 Hs.145998	ESTs, Weakly similar to PN0099 son3 prot ESTs	23.20		2.10
50	106507	AA259068	Hs.267819	protein phosphatase 1, regulatory (inhib	15.20		
	106490	AA404265	Hs.115537	putative dipeptidese			
	106474	BE383668	Hs.42484	hypothetical protein FLJ10618	10.44		
~ ~	106211	AA428240	Hs.126083	ESTs		29.80	
55	105986	AB037722	Hs.8707	KIAA1301 protein		3.70	4.04
	105894	A1904740	Hs.25691	receptor (calcitonin) activity modifying			1.94 1.75
	105847	AW964490 AW747996	Hs.32241	ESTs, Weakly similar to S65657 alpha-1C- ESTs, Moderately similar to A56194 throm			2.47
	105803 105731	AA834664	Hs.160999 Hs.29131	nuclear receptor coactivator 2	10.71	•	
60	105731	H46612	Hs.293815	Homo sapiens HSPC285 mRNA, partial cds			
•	105688	Al299139	Hs.17517	ESTs	23.40		
	105510	Z42047	Hs.283978	Homo sapiens PRO2751 mRNA, complete cds	37.20	0.00	
	105101	H63202	Hs.38163	ESTs		8.30	
65	104989	R65998	Hs.285243	hypothetical protein FLJ22029		8.09	1.92
65	104986	AW088826	Hs.117176	poly(A)-binding protein, nuclear 1 phosphatidylinositol-4-phosphate 5-kinas		5.40	1.52
	104969 104903	A1670947 A1436323	Hs.78406 Hs.31141	Homo saplens mRNA for KIAA1568 protein,		7.60	
	104896	AW015318	Hs.23165	ESTs	13.80		
	104865	T79340	Hs.22575	Homo saplens cDNA: FLJ21042 fis, clone C			
70	104825	AA035613	Hs.141883	ESTs			1.87
	104781	AA099904	Hs.21610	DKFZP434B203 protein			1.93
	104776	AA026349		gb:zj99f01.s1 Soares_pregnant_uterus_NbH		10.20	
	104691	U29690	Hs.37744	Homo saplens beta-1 adrenergic receptor		5.69	
75	104667	AI239923	Hs.30098	eSTs gb:EST00057 HE6W Homo saplens cDNA clone		3.82 4.20	
13	104404 104392	H58762 AA076049	Hs.274415	Homo saplens cDNA FLJ10229 fis, clone HE	27.20	7.20	
	104392	AB002298	Hs.173035	KIAA0300 protein	** ***		1.91
	104074	AL162039	Hs.31422	Homo saplens mRNA; cDNA DKFZp434M229 (fr	11.20		
	103749	AL135301	Hs.8768	hypothetical protein Fلــا10849	10.86		
80	103645	AW246253	Hs.7043	succinate-CoA ligase, GDP-forming, alpha	12.00		4
	103554	A1878826	Hs.323469	caveolin 1, caveolae protein, 22kD			- 1.80
	103541	AI815601	Hs.79197	CD83 antigen (activated B lymphocytes, i			
	103496	Y09267	Hs.132821	flavin containing monooxygenase 2 A kinase (PRKA) anchor protein 1	11.20		
85	103428 103353	BE383507 X89399	Hs.78921 Hs.119274	RAS p21 protein activator (GTPase activa	19.80		
00	100000	AUJUJJ	119.113214	10 to he : higher content for the nonte	.5.00		

WO 02/086443								PCT/US02/12476
	103295	X81479	Hs.2375	eaf-like module containing, mucin-like,		3.60		
	103280	U84722	Hs.76206	cadherin 5, type 2, VE-cadherin (vascula				
	103100	NM_005574	Hs.184585	LIM domain only 2 (rhombotin-like 1)			1.76	
	103025	NM_002837	Hs.123641	protein tyrosine phosphatase, receptor t			2.15	
5	102698	M18667	Hs.1867	progastricsin (pepsinogen C)				
	102659	BE245169	Hs.211610	CUG triplet repeat, RNA-binding protein	11.00			
	102580	U60808	Hs.152981	CDP-diacyiglycerol synthase (phosphatida	25.40			
	102417	AA034127	Hs.153487	signal transducing adaptor molecule (SH3	14.00			•
	102363	NM_003734	Hs.198241	amine oxidase, copper containing 3 (vasc				
10		AA306342	Hs.69171	protein kinase C-like 2	10.86			
	102283	AW161552	Hs.83381	guanine nucleotide binding protein 11				
	102188	U20350	Hs.78913	chemokine (C-X3-C) receptor 1		7.40		
	102151	T27013	Hs.3132	steroidogenic acute regulatory protein	16.40			
4 ~	101957	L28824	Hs.74101	spleen tyrosine kinase	15.40			
15	101842	M93221	Hs.75182	mannose receptor, C type 1	•			
	101771	NM_002432	Hs.153837	myeloid cell nuclear differentiation ant			1.78	
		Al198550	Hs.81256	S100 calcium-binding protein A4 (calcium	40.00		1.78	
	101716	AF050658	Hs.2563	tachykinin, precursor 1 (substance K, su	18.80		2.22	
20	101678	M62505	Hs.2161	complement component 5 receptor 1 (C5a)	504.80		2.22	
20	101447	M21305	11	gb:Human alpha satellite and satellite 3	304.60	31.00		
	101383	NM_000132	Hs.79345	coagulation factor VIII, procoagulant co		31.00	1.75	
	101346	Al738616	Hs.77348	hydroxyprostaglandin dehydrogenase 15-(N			1.75	
	101345		Hs.152175 Hs.75678	calcitonin receptor-like FBJ murine osteosarcoma viral oncogene h			2.24	
25	101336 101330	NM_006732 L43821	Hs.80261	enhancer of filamentation 1 (cas-like do			Ga 4.7	
23		BE297626	Hs.296049	microfibrillar-associated protein 4				
	101277 101262		NS.230043	gb:Human dystrophin (dp140) mRNA, 5' end	19.00			
	101168	NM_005308	Hs.211569	G protein-coupled receptor kinase 5			2.01	
	101102	NM_003243	Hs.79059	transforming growth factor, beta recepto				
30	101088	X70697	Hs.553	solute carrier family 6 (neurotransmitte		7.52		
50	101066	AW970254	Hs.889	Charot-Leyden crystal protein	19.38	***		
	100971	BE379727	Hs.83213	fatty acid binding protein 4, adipocyte			1.91	
	100893	BE245294	Hs.180789	S164 protein	15.40			
	100770	W25797.comp		amyloid beta (A4) precursor protein (pro	11.20			
35	100716	X89887	Hs.172350	HIR (histone cell cycle regulation defec	14.80			
	100555	M69181		gb:Human nonmuscle myosin heavy chain-B	33.00			
	100425	NM_014747	Hs.78748	KIAA0237 gene product	16.20			
	100408	D86640	Hs.56045	src homology three (SH3) and cysteine ri		4.00		
40	100382	D83407	Hs.156007	Down syndrome critical region gene 1-lik		4.24		
40	100351	D64158				6.20		
	100299	D49493	Hs.2171	growth differentiation factor 10		21.20		
		AA305746	Hs.49	macrophage scavenger receptor 1			1.79	
	100108	U09577	Hs.76873	hyaluronoglucosaminidase 2		5.40	1.78	
15	100095	Z97171	Hs.78454	myodlin, trabecular meshwork inducible	11.29	3.40		
45	100066			•	11.23			

TABLE 3B shows the accession numbers for those primekeys tacking unigenelD's for Table 3A. For each probeset we have listed the gene cluster number from which the oligonucleotides were designed. Gene clusters were compiled using sequences derived from Genbank ESTs and mRNAs. These sequences were clustered based on sequence similarity using Clustering and Alignment Tools (DoubleTwist, Oakland California). The Genbank accession numbers for sequences comprising each cluster are listed in the "Accession" column.

Pkey: Unique Eos probeset identifier number CAT number: Gene cluster number Accession: Genbank accession numbers

60	Pkey	CAT number	Accessions	
00	123619	371681_1	AA602964 A	A609200
	126433		AA325606 A	A099517 N89423
	125831	1522905 1		
	126816	122973_1	AA248234 A	
65	126852	136135 1	AA399961 A	
•	121059			
	120637			A809404 AA286907 AW977624
	122011			
	120934			A226513 AA383773
70	123802			AA620448
	116814			H50834
	118329			N63520
	104404		H58762	
	104776			AA026349
75	113502			
	101262			
	108573			AA086005
	101447			, 1,100000
	124357			N22401
80	108781			AA128654
00	112794	genbank_R9701		R97018
	100351			101010
	100555	tigr_HT2245		1105 1151039
	100000	081_1112240	141031011100	1,00 001000

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Table 4A shows 202 genes up-regulated in samples from patients treated with chemotherapy or radiotherapy. These genes were selected from 59680 probesets on the Eos/Affymetrix Hu03 Genechip array. Gene expression data for each probeset obtained from this analysis was expressed as average Intensity (AI), a normalized value reflecting the relative level of mRNA expression.

Pkey: Unique Eos probeset identifier number

ExAccn: Exemplar Accession number, Genbank accession number

UnigenelD: Unigene number

Unigene Title: Unigene gene title

R1: average of Al for samples from patients treated with chemotherapy or radiotherapy divided by the average of Al for normal lung samples. 5

10	R1:	average of	Al for samples	s from patients treated with chemotherapy or radiother	apy divided
10	Pkey	ExAcon	UnigenelD	Unigene Title	R1
	,	2.0 .00	·		
	100113	NM_001269	Hs.84746	chromosome condensation 1	27.20
1.5	100187	D17793	Hs.78183	aldo-keto reductase family 1, member C3	20.60
15	100210	D26361	Hs.3104	KIAA0042 gene product glutamate receptor, metabotropic 5	20.40 20.60
	100225 100269	D28539 NM_001949	Hs.167185 Hs.1189	E2F transcription factor 3	29.40
	100438	AA013051	Hs.91417	topolsomerase (DNA) II binding protein	23.50
	100877	X80821	Hs.27973	KIAA0874 protein	35.56
20	100893	BE245294	Hs.180789	S164 protein	43.40
	101273	Z11933	Hs.182505	POU domain, class 3, transcription facto	21.80
	101447	M21305	H- 4600	gb:Human alpha satellite and satellite 3	193.60 38.40
	101649 101724	AW959908 L11690	Hs.1690 Hs.620	heparin-binding growth factor binding pr bullous pemphigoid antigen 1 (230/240kD)	198.80
25	101748	NM_001944	Hs.1925	desmoglein 3 (pemphigus vulgaris antigen	78.60
	101809	M86849	Hs.323733	gap junction protein, beta 2, 26kD (conn	162.20
	101879	AA176374	Hs.243886	nuclear autoantigenic sperm protein (his	50.00
	101915	AF207881	Hs.155185	cytosolic ovarian carcinoma antigen 1	26.00
20	101973	U41514	Hs.80120	UDP-N-acetyl-alpha-D-galactosamine:polyp	37.20
30	102025 102031	U04045 U04898	Hs.78934 Hs.2156	mutS (E. coli) homolog 2 (colon cancer, RAR-related orphan receptor A	32.00
	102052	NM_002202	Hs.505	ISL1 transcription factor, LIM/homeodoma	51.20
	102391	AA296874	Hs.77494	deoxyguanosine kinase	13.90
	102420	U44060	Hs.14427	Homo sapiens cDNA: FLJ21800 fis, clone H	28.80
35	102610	U65011	Hs.30743	preferentially expressed antigen in mela	110.60
	102829	NM_006183	Hs.80962	neurotensin	116.80 2.30
	103000	NM_001975	Hs.146580 Hs.83169	enolase 2, (gamma, neuronal) matrix metalloprotelnase 1 (Interstitial	181.40
	103036 103507	M13509 AJ000512	Hs.296323	serum/glucocorticoid regulated kinase	49.20
40	103587	BE270266	Hs.82128	5T4 oncofetat trophoblast glycoprotein	86.60
	104660	BE298665	Hs.14846	Homo sapiens mRNA; cDNA DKFZp564D016 (fr	42.60
	104896	AW015318	Hs.23165	ESTs	29.40
	105038	AW503733	Hs.9414	KIAA1488 protein	21.50 32.80
45	105298	BE387790	Hs.26369 Hs.283978	hypothetical protein FLJ20287 Homo saplens PRO2751 mRNA, complete cds	20.20
73	105510 105667	Z42047 AA767526	Hs.22030	paired box gene 5 (B-cell lineage specif	28.40
	106073	AL157441	Hs.17834	downstream neighbor of SON	25.40
	106205	AW965058	Hs.111583	ESTs, Weakly similar to I38022 hypotheti	32.00
50	106516	AL137311	Hs.234074	Homo sapiens mRNA; cDNA DKFZp761G02121 (40.60
50	106533	AL134708	Hs.145998	ESTs ESTs	59.80 43.40
	106575 106654	AW970602 AW075485	Hs.105421 Hs.286049	phosphoserine aminotransferase	50.80
	106851	A)458623	113.2000-10	gb:tk04g09.x1 NCI_CGAP_Lu24 Homo saplens	53.40
	106995	AB023139	Hs.37892	KIAA0922 protein	20.88
55	107332	T87750	Hs.183297	DKFZP566F2124 protein	23.60
	107532	AA443473	Hs.173684	Homo sapiens mRNA; cDNA DKFZp762G207 (fr	57.20
	107922	BE153855	Hs.61460 Hs.69499	lg superfamily receptor LNIR hypothetical protein	49.00 19.67
	108609 108780	BE409857 AU076442	Hs.117938	collagen, type XVII, alpha 1	48.17
60	109166	AA219691	Hs.73625	RAB6 interacting, kinesin-like (rabkines	59.20
	109260	AW978515	Hs.131915	KIAA0863 protein	28.60
	109280	AK001355	Hs.279610	hypothetical protein FLJ10493	22.80
	109292	AW975746	Hs.188662	KIAA1702 protein ESTs	21.00
65	109384 109415	AA219172 U80736	Hs.86849 Hs.110826	trinucleotide repeat containing 9	31.60
UJ	109445	AA232103	Hs.189915	ESTs	24.20
	109502	AW967069	Hs.211556	hypothetical protein MGC5487	21.40
	109633	AW003785	Hs.170267	ESTs	20.40
70	109786	AI989482	Hs.146286	kinesin family member 13A	19.60
70 ·	109958	AA001266	Hs.133521	ESTs HMT1 (hnRNP methyltransferase, S. cerevi	24.00 28.40
	110920 110924	N47224 AW058463	Hs.20521 Hs.12940	zinc-fingers and homeoboxes 1	36.00
	111084	H44186	Hs.15456	PDZ domain containing 1	61.20
	111132	AB037807	Hs.83293	hypothetical protein	24.60
75	111229	AW389845	Hs.110855	ESTs	27.20
	111337	AA837396	Hs.263925	LIS1-interacting protein NUDE1, rat homo	48.00
	111987	NM_015310	Hs.6763 Hs.22116	KIAA0942 protein CDC14 (cell division cycle 14, S. cerevi	37.80 26.80
	112046 112268	AA383343 W39609	Hs.22116 Hs.22003	solute carrier family 6 (neurotransmitte	63.80
80	112685	R87650	Hs.33439	ESTs, Weakly similar to ALU1_HUMAN ALU	26.40
	112871	AL110216	Hs.12285	ESTs, Weakly similar to !55214 salivary	47.64
	112897	AW206453	Hs.3782	ESTs	22.00
	112973	AB033023	Hs.318127	hypothetical protein FLJ10201	65.00
85	112992 113073	AL157425 N39342	Hs.133315 Hs.103042	Homo sapiens mRNA; cDNA DKFZp761J1324 (f microtubule-associated protein 1B	42.00 55.40
05	113013	1700074	. 10.100072		

	W	O 02/0864	143		
	113494	T91451	Hs.86538	ESTs	22.80
	113560	T91015	Hs.268626	ESTs	22.80
	113849	AA457211	Hs.8858	bromodomain adjacent to zinc finger doma	51.80
5	113950	AI267652	Hs.30504	Homo sapiens mRNA; cDNA DKFZp434E082 (fr	28.20
J	114339	AA782845	Hs.22790	ESTs	20.20 21.00
	114365 114455	H42169 H37908	Hs.18653 Hs.271616	hypothetical protein FLJ14627 ESTs, Weakly similar to ALUB_HUMAN ALU S	25.80
	114518	AW163267	Hs.106469	suppressor of var1 (S.cerevislae) 3-like	23.60
	114824	AA960961	Hs.305953	zinc finger protein 83 (HPF1)	27.20
10	114837	BE244930	Hs.166895	ESTs	30.20
	114974	AW966931	Hs.179662	nucleosome assembly protein 1-like 1	20.80
•	115075	AA814043	Hs.88045	ESTs	30.60
	115084	BE383668	Hs.42484	hypothetical protein FLJ10618	28.86
15	115291	BE545072	Hs.122579	hypothetical protein FLJ10461	38.00 22.60
13	115313	AA808001 D31382	Hs.184411 Hs.63325	albumin transmembrane protease, serine 4	173.60
	115697 115909	AW872527	Hs.59761	ESTs, Weakly similar to DAP1_HUMAN DEATH	27.77
	116090	AI591147	Hs.61232	ESTS	20.80
	116107	AL133916	Hs.172572	hypothetical protein FLJ20093	164.20
20	116399	AA889120	Hs.110637	homeo box A10	38.00
	117099	H93699		gb:yv16a11.s1 Soares fetal liver spleen	21.60
	117881	AF161470	Hs.260622	butyrate-induced transcript 1	49.40
	118091	AW005054	Hs.47883	ESTs, Weakly similar to KCC1_HUMAN CALCI	22.40 22.00
25	118138 118720	AA374756 N73515	Hs.93560	Homo sapiens mRNA for KIAA1771 protein, gb:za49d07.s1 Soares fetal liver spleen	20.00
	118873	A1824009	Hs.44577	ESTs	19.40
	119126	R45175	Hs.117183	ESTs	111.20
	119717	AA918317	Hs.57987	B-cell CLL/lymphoma 118 (zinc finger pro	33.00
20	119940	AL050097	Hs.272531	DKFZP586B0319 protein	31.00
30	120266	AI807264	Hs.205442	ESTs, Weakly similar to T34036 hypotheti	20.20
	120515	AA258356) le 4040	gb:zr59c10.s1 Soares_NhHMPu_S1 Homo sapi achaete-scute complex (Drosophila) homol	25.00 95.40
	120859 120983	AA826434 AA398209	Hs.1619 Hs.97587	EST	105.20
	121054	AW976570	Hs.97387	ESTs	38.80
35	121369	AW450737	Hs.128791	CGI-09 protein	41.60
	122335	AA443258	Hs.241551	chloride channel, calcium activated, fam	30.80
	122612	AA974832	Hs.128708	ESTs	19.60
	123130	AA487200		gb:ab19f02.s1 Stratagene lung (937210) H	33.20
40	123440	A1733692	Hs.112488	ESTs	23.17
40	123596 123619	AA421130 AA602964	Hs.112640	EST gb:no97c02.s1 NCI_CGAP_Pr2 Homo sapiens	23.00 28.80
	124006	Al147155	Hs.270016	ESTs .	77.60
	124169	BE079334	Hs.271630	ESTs	22.20
	124281	Al333756	Hs.111801	arsenate resistance protein ARS2	42.20
45	124472	N52517	Hs.102670	EST	32.60
	124617	AW628168	Hs.152684	ESTs	21.80
	124631	NM_014053	Hs.270594	FLVCR protein	30.40 21.20
	124839 125186	R55784 AA610620	Hs.140942 Hs.181244	ESTs major histocompatibility complex, class	42.80
50	125321	T86652	Hs.178294	ESTs	27.00
	125535	NM_013243	Hs.22215	secretogranin III	23.80
	125646	AA628962	Hs.75209	protein kinase (cAMP-dependent, catalyti	23.20
	125684	AW589427	Hs.158849	Homo saplens cDNA: FLJ21663 fis, clone C	21.20
55	125724	AL360190	Hs.295978	Homo sapiens mRNA full length insert cDN	48.80
23	125847 125934	AW161885 AA193325	Hs.249034 * Hs.32646	ESTs hypothetical protein FLJ21901	31.00 21.20
	126077	M78772	Hs.210836	ESTs	49.80
	126299	AW979155	Hs.298275	amino acid transporter 2	21.80
	126395	A1468004	Hs.278956	hypothetical protein FLJ12929	71.00
60	126433	AA325606		gb:EST28707 Cereballum II Homo saplens c	23.20
	126509	R47400	Hs.23850	ESTs	23.80
	126538	AB030656	Hs.17377	coronin, actin-binding protein, 1C	23.10 36.00
	126666 126812	AA648886 AB037860	Hs.151999 Hs.173933	ESTs nuclear factor I/A	20.80
65	126872	AW450979	110.170000	gb:UI-H-BI3-ala-a-12-0-UI.s1 NCI_CGAP_Su	46.29
	127046	AA321948	Hs.293968	ESTs	22.80
	127431	AW771958	Hs.175437	ESTs, Moderately similar to PC4259 ferri	30.00
	127489	AA650250	Hs.272076	ESTs	20.80
70	127521	AW297206	Hs.164018	ESTs	25.20
70	127742	AW293496 AA805151	Hs.180138	ESTs mitogen-activated protein kinase kinase	28.00 21.20
	127925 127930	AA809672	Hs,3628 Hs.123304	ESTs	20.54
	127968	AA830201	Hs.124347	ESTs	28.20
	127987	Al022103	Hs.124511	ESTs	19.60
75	128116	H07103	Hs.286014	Homo sapiens, clone IMAGE:3867243, mRNA	20.40
	128609	NM_003616	Hs.102456	survival of motor neuron protein interac	34.40
	128777	Al878918	Hs.10526	cysteine and glycine-rich protein 2	53.80
	128949	AA009647	Hs.8850	a disintegrin and metalloproteinase doma	23.00
80	129168	Al132988 Al267700	Hs.109052 Hs.317584	chromosome 14 open reading frame 2 ESTs	37.60 28.60
30	129404 129527	AA769221	Hs.270847	delta-tubulin .	40.80
	129574	AA026815	Hs.11463	UMP-CMP kinase	31.20
	129598	N30436	Hs.11556	Homo sapiens cDNA FLJ12566 fis, clone NT	29.60
05	129785	H19006	Hs.184780	ESTs	72.20
85	129970	AV655806	Hs.296198	chromosome 12 open reading frame 4	22.20

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	**	0 02,000			
	130149	AW067805	Hs.172665	methylenetetrahydrofolate dehydrogenase	29.60
	130199	Z48579	Hs.172028	a disintegrin and metalloproteinase doma	27.60
	130441	U63630	Hs.155637	protein kinase, DNA-activated, catalytic	28.36
_	130466	W19744	Hs.180059	Homo sapiens cDNA FLJ20653 fis, clone KA	20.20
5	130482	AW409701	Hs.1578	baculoviral IAP repeat-containing 5 (sur	22.40
	130617	M90516	Hs.1674	glutamine-fructose-6-phosphate transamin	19.60
	130703	R77776	Hs.18103	ESTs	19.40
	130732	AW890487	Hs.63984	cadherin 13, H-cadherin (heart)	21.40
	130867	NM_001072	Hs.284239	UDP glycosyltransferase 1 family, polype	110.00
10	131028	Al879165	Hs.2227	CCAAT/enhancer binding protein (C/EBP),	25.20
	131086	AL035461	Hs.2281	chromogranin B (secretogranin 1)	40.60
	131284	NM_001429	Hs.25272	E1A binding protein p300	24.60
	131775	AB014548	Hs.31921	KIAA0648 protein	21.00
	131860	BE383676	Hs.334	Rho quanine nucleotide exchange factor (33.40
15	131945	NM_002916	Hs.35120	replication factor C (activator 1) 4 (37	60.80
15	132040	NM_001196	Hs.315689	Homo saplens cDNA: FLJ22373 fis, clone H	20.40
	132084	NM_002267	Hs.3886	karyopherin alpha 3 (importin alpha 4)	29.40
	132389	AA310393	Hs.190044	ESTs	32.40
	132437	AA152106	Hs.4859	cyclin L ania-6a	27.40
20	132550	AW969253	Hs.170195	bane morphogenetic protein 7 (osteogenic	75.60
20	132617	AF037335	Hs.5338	carbonic anhydrase XII	31.36
	132632	AU076916	Hs.5398	quanine monphosphate synthetase	32.40
	132672	W27721	Hs.54697	Cdc42 quanine exchange factor (GEF) 9	23,40
	132742	AA025480	Hs.292812	ESTs, Weakly similar to T33468 hypotheti	61.20
25	132771	Y10275	Hs.56407	phosphoserine phosphatase	22.33
23	133070	U92649	Hs.64311	a disintegrin and metalloproteinase doma	23.50
	133153	AF070592	Hs.66170	HSKM-B protein	30.00
	133181	X91662	Hs.66744	twist (Drosophila) homolog (acrocephalos	23.80
	133282	AA449015	Hs.286145	SRB7 (suppressor of RNA polymerase B, ye	51.60
30	133350	AI499220	Hs.71573	hypothetical protein FLJ10074	33.00
20	133592	AV652066	Hs.75113	general transcription factor IIIA	82.00
	133658	AV052005 AA319146	Hs.75426	secretogranin II (chromogranin C)	02.00
			Hs.170290	discs, large (Drosophila) homolog 5	69.33
	133865	AB011155 NM 005025	Hs.78589	serine (or cystelne) proteinase inhibito	33.20
35	134032		Hs.50421	KIAA0203 gene product	31.60
33	134125	NM_014781	Hs.79428	BCL2/adenovirus E1B 19kD-interacting pro	30.60
	134158	U15174 BE538082	Hs.8172	ESTs, Moderately similar to A48010 X-lin	23.40
	134321		Hs.82285	phosphoribosylglycinemide formyltransfer	49.20
	134367	AA339449 U66615	Hs.172280	SWI/SNF related, matrix associated, acti	20.20
40	134570		Hs.173135	dual-specificity tyrosine-(Y)-phosphoryl	20.80
40	134753	NM_006482 AA448542	Hs.251677	Gantigen 7B	37.60
	135002			hydroxysteroid (17-beta) dehydrogenase	53.40
	135029	H58818	Hs.187579	cyclin-dependent kinase 5, regulatory su	31.60
	135047	AL134197	Hs.93597 Hs.99171	neurotrophin 3	28.80
45	135345	X53655	ו זו פט.פה	neurosopian o	20.00
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TABLE 4B shows the accession numbers for those primekeys lacking unigenelD's for Teble 4A. For each probeset we have listed the gene cluster number from which the oligonucleotides were designed. Gene clusters were compiled using sequences derived from Genbank ESTs and mRNAs. These sequences were clustered based on sequence similarity using Clustering and Alignment Tools (DoubleTwist, Oakland California). The Genbank accession numbers for sequences comprising each cluster are listed in the

Pkey: Unique Eos probeset identifier number CAT number: Gene cluster number Accession: Genbank accession numbers

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	Pkey	CAT number	Accessions
	123619	371681_1	AA602964 AA609200
60	126433	127143_1	AA325606 AA099517 N89423
	126872	142696 1	AW450979 AA136653 AA136656 AW419381 AA984358 AA492073 BE168945 AA809054 AW238038 BE011212 BE011359
			BE011367 BE011368 BE011362 BE011215 BE011365 BE011363
	106851	322947_1	Al458623 AA639708 AA485409 R22065 AA485570
	118720	genbank_N73518	6 N73515
65	120515	genbank_AA258	356 AA258356
	117099		H93699 H97976 H80036
	101447	entrez_M21305	M21305
	123130	genhank AA487	200 AA487200

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Table 5A shows 680 genes up-regulated in squamous cell carcinoma or adenocarcinoma lung tumors relative to normal lung and chronically diseased lung. These genes were selected from 59680 probesets on the Eos/Affymetrix Hu03 Genechip array. Gene expression data for each probeset obtained from this analysis was expressed as average intensity (AI), a normalized value reflecting the relative level of mRNA expression.

5 Pkey: Unique Eos probeset identifier number Exemplar Accession number, Genbank accession number ExAccn: UnigenelD: Unigene Title: R1: Unigene number Unigene gene title 70th percentile of Al for squamous cell carcinoma and adenocarcinoma lung tumor samples divided by the 90th percentile of Al for normal and chronically diseased lung samples.

80th percentile of AI adenocarcinoma lung tumor samples divided by the 90th percentile of AI for normal and chronically diseased lung samples.

80th percentile of AI guamous cell carcinoma lung tumor samples divided by the 90th percentile of AI for normal and chronically diseased lung samples.

80th percentile of AI adenocarcinoma lung tumor samples divided by the 80th percentile of AI for squamous cell carcinoma lung tumor samples.

70th percentile of AI for squamous cell carcinoma and adenocarcinoma lung tumor samples minus the 15th percentile of AI for all normal lung, chronically diseased lung and tumor samples divided by 90th percentile of AI for normal and chronically diseased lung samples minus the 15th percentile of AI for all normal lung, chronically diseased lung and tumor samples 10 R2: R3: R4:

R5: 15

20	Pkey	ExAcon	UnigenelD	Unigene Title	R1	R2	R3	R4	R5
20	100035			AFFX control: GAPDH					6.76
	100036			AFFX control: GAPDH					5.77
	100037	,		AFFX control: GAPDH		8.00			5.75
25	100071	A28102 X02308	Hs.82962	Human GABAa receptor alpha-3 subunit thymidylate synthetase		0.00			5.71
23	100114 100154	H60720	Hs.81892	KIAA0101 gene product	3.84				
	100187	D17793	Hs.78183	aldo-keto reductase family 1, member C3	3.33				
	100188	AW247090	Hs.57101	minichromosome maintenance deficient (S.					4.52
20	100202	BE294407	Hs.99910	phosphofructokinase, platelet					5.49 5.67
30	100216 100269	AA489908 NM_001949	Hs.1390 Hs.1189	proteasome (prosome, macropain) subunit, E2F transcription factor 3	2.55				0.0.
	100203	AU076657	Hs.1600	chaperonin containing TCP1, subunit 5 (e					5.66
	100297	AU077258	Hs.182429	protein disulfide isomerase-related prot					3.81
25	100330	AW410976	Hs.77152	minichromosome maintenance deficient (S.	· 5.07				4.50
35	100335 100360	AW247529	Hs.6793 Hs.75939	platelet-activaling factor acetylhydrola uridine monophosphate kinase	5.07				4.82
	100300	W70171 NM_014791	Hs.184339	KIAA0175 gene product					3.79
	100474	NM_000699	Hs.300280	amylase, alpha 2A; pancreatic				15.65	
40	100486	T19006	Hs.10842	RAN, member RAS oncogene family					5.49
40	100491	D56165	Hs.275163	non-metastatic cells 2, protein (NM23B)		7.20			4.17
	100516	D90278	Hs.11 Hs.99949	carcinoembryonic antigen-related cell ad protectin-induced protein		1.20		14.20	
	100522 100559	X51501 NM_000094	Hs.1640	collagen, type VII, alpha 1 (epidermolys	3.10				
	100576	X00356	Hs.37058	calcitonin/calcitonin-related polypeptid				9.30	
45	100629	AA015693	Hs.21291	mitogen-activated protein kinase kinase				20.60	
	100661	BE623001	Hs.132748	Homo sapiens ribosomal protein L39 mRNA,	3.85	8.60			
	100677 100696	AA353686 D14887	Hs.57813 Hs.121686	zinc ribbon domain containing, 1 general transcription factor IIA, 1 (37k		0.00		10.00	
	100709	N26539	Hs.100469	myeloid/lymphold or mixed-lineage leukem			24.80		
50	100761	BE208491	Hs.295112	KIAA0618 gene product		7.60			
	100830	AC004770	Hs.4756	flap structure-specific endonuclease 1		40.00			7.99
	100867	U14622	LIA 207270	gb:Human transketolase-like protein gene ret proto-oncogene (multiple endocrine n		10.20 8.00			
	100902 100906	M16029 AU076916	Hs.287270 Hs.5398	quanine monphosphale synthetase		0.00			5.16
55	100960	J00124	Hs,117729	keratin 14 (epidermolysis bullosa simple	2.57				
	101045	J05614		gb:Human proliferating cell nuclear anti					4.69
	101061	NM_000175	Hs.180532	glucose phosphate isomerase		12.91			4.19
	101071. 101124	L02840 L10343	Hs.84244 Hs.112341	potassium voltage-gated channel, Shab-re protease inhibitor 3, skin-derived (SKAL	3.12	12.51			
60	101175	U82671	Hs.36980	melanoma antigen, family A, 2	3.50				
•	101181	BE262621	Hs.73798	macrophage migration inhibitory factor (5.69
	101204	L24203	Hs.82237	ataxia-telangiectasia group D-associated	4.08		6.40		
	101210	L29301	Hs.2353	opioid receptor, mu 1 cyclin-dependent kinase inhibitor 3 (CDK	2.53		0.40		
65	101216 101228	AA284166 AA333387	Hs.84113 Hs.82916	chaperonin containing TCP1, subunit 6A (2.00				7.90
05	101233	AL135173	Hs.878	sorbitol dehydrogenase					4.45
	101273	Z11933	Hs.182505	POU domain, class 3, transcription facto	8.50				4.47
	101342	U52112	Hs.182018	Interleukin-1 receptor-associated kinase				21.89	4.17
70	101346 101369	AI738616 NM_000892	Hs.77348 Hs.1901	hydroxyprostaglandin dehydrogenase 15-(N kaliikrein B, plasma (Fletcher factor) 1				12.80	
70	101396	BE267931	Hs.78996	proliferating cell nuclear antigen	3.24				
	101431	BE185289	Hs.1076	small proline-rich protein 1B (comifin)					7.90
	101448	NM_000424	Hs.195850	keratin 5 (epidermolysis bullosa simplex	8.31			38.80	
75	101462	AL035668	Hs.73853	bone morphogenetic protein 2 glutamic-oxaloacetic transaminase 2, mit				35.00	4.01
13	101466 101484	BE262660 AA053486	Hs.170197 Hs.20315	interferon-induced protein with tetratri				12.00	7.01
	101502	M26958	110,20010	gb:Human parathyroid hormone-related pro	10.50				
	101505	AA307680	Hs.75692	asparagine synthetase					4.46
οΛ	101526	NM_002197	Hs.154721	aconitase 1, soluble	4.02				4.65
80	101535 101577	X57152 M34353	Hs.99853 Hs.1041	fibrillarin v-ros avian UR2 sarcoma virus oncogene h				9.09	7.00
	101649	AW959908	Hs.1690	heparin-binding growth factor binding pr	54.00				
	101663	NM_003528	Hs.2178	H2B histone family, member Q	5.59				
O.F	101664	AA436989	Hs.121017	H2A histone family, member A	7.00	7.00			
85	101669	L24498	Hs.80409	growth arrest and DNA-damage-inducible,		7.60			

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	101695	M69136	Hs.135626	chymase 1, mast cell	4.79				
	101724	L11690	Hs.620	bullous pemphigold antigen 1 (230/240kD)	15.21				
	101748	NM_001944	Hs.1925	desmoglein 3 (pemphigus vulgaris antigen	55.50				4.10
5	101759 101771	M80244 NM_002432	Hs.184601 Hs.153837	solute carrier family 7 (cationic amino myeloid cell nuclear differentiation ant				18.57	4.10
,	101804	M86699	Hs.169840	TTK protein kinase	4.50				
	101809	M86849	Hs.323733	gap junction protein, beta 2, 26kD (conn	140.00				
	101833	AU076442	Hs.117938	collagen, type XVII, alpha 1	2.56				
10	101842	M93221	Hs.75182	mannose receptor, C type 1				12.80	c 00
10	101851	BE260964	Hs.82045	midkine (neurite growth-promoting factor		7.80			5.88
	102002 102039	NM_002484 AL134223	Hs.81469 Hs.306098	nucleotide binding protein 1 (E.coll Min aldo-keto reductase family 1, member C1		7.00			4.35
	102072	U09410	Hs.78743	zinc finger protein 131 (clone pHZ-10)			7.40		
	102083	T35901	Hs.75117	interleukin enhancer binding factor 2, 4					5.12
15	102111	L36196	Hs.81884	sulfotransferase family, cytosolic, 2A,				12.00	
	102123	NM_001809	Hs.1594	centromere protein A (17kD)	6.20 2.62				
	102154 102193	U17760 AL036335	Hs.75517 Hs.313	taminin, beta 3 (niceln (125kD), kalinin secreted phosphoprotein 1 (osteopontin,	5.85				
	102217	AA829978	Hs.301613	JTV1 gene	0.00				6.18
20	102224	NM_002810	Hs.148495	proteasome (prosome, macropain) 26S subu					4.49
	102234	AW163390	Hs.278554	heterochromatin-like protein 1					5.80
	102251	NM_004398	Hs.41706	DEAD/H (Asp-Glu-Ala-Asp/His) box polypep	4.50				5.15
	102305 102330	AL043202 BE298063	Hs.90073 Hs.77254	chromosome segregation 1 (yeast homolog) chromobox homolog 1 (Drosophila HP1 beta					4.17
25	102340	U37055	Hs.278657	macrophage stimulating 1 (hepatocyte gro				9.33	
	102348	U37519	Hs.87539	aldehyde dehydrogenase 3 family, member	8.87				
	102368	U39817	Hs.36820	Bloom syndrome	15.91		40.00		
	102394	NM_003816	Hs.2442	a disintagrin and metalloproteinase doma vascular endothelial growth factor C			19.20	14.00	
30	102404 102537	NM_005429 U57094	Hs.79141 Hs.50477	RAB27A, member RAS oncogene family				12.00	
30	102581	AU077228	Hs.77256	enhancer of zeste (Drosophila) homolog 2					4.57
	102605	Al435128	Hs.181369	ubiquitin fusion degradation 1-like					3.98
	102610	U65011	Hs.30743	preferentially expressed antigen in mela	77.50				
35	102623	AW249285	Hs.37110 Hs.23016	melanoma antigen, family A, 9 G protein-coupled receptor	12,50		22.00		
23	102642 102654	AA205847 AV649989	Hs.24385	Human hbc647 mRNA sequence		12.00	22.00		
	102659	BE245169	Hs.211610	CUG triplet repeat, RNA-binding protein				12.80	
	102669	U71207	Hs.29279	eyes absent (Drosophila) homolog 2	6.50				
40	102672	U72066	Hs.29287	retinoblastoma-binding protein 8	8.50				9.24
40	102687 102696	NM_007019 BE540274	Hs.93002 Hs.239	ubiquitin carrier protein E2-C forkhead box M1					5.54
	102768	U82321	. 15.250	gb:Homo sapiens clone 14.98 mRNA sequenc		6.60			
	102781	BE258778	Hs.108809	chaperonin containing TCP1, subunit 7 (e					3.78
45	102784	U8565B	Hs.61796	transcription factor AP-2 gamma (activat			14.40		4.26
40	102824 102829	U90916 NM_006183	Hs.82845 Hs.80962	Homo sapiens cDNA: FLJ21930 fis, clone H neurotensin	8.00		14.40		
	102888	Al346201	Hs.76118	ubiquilin carboxyl-terminal esterase L1	5.55				5.50
	102892	BE440042	Hs.83326	matrix metalloproteinase 3 (stromelysin			6.70		
50	102913	NM_002275	Hs.80342	keratin 15	4.64				
50	102935 102951	BE561850 X15218	Hs.80506 Hs.2969	smail nuclear ribonucleoprolein polypept v-ski avian sarcoma viral oncogene homol	2.93			11.40	
	102983	BE387202	Hs.118638	non-metastatic cells 1, protein (NM23A)					7,26
	103023	AW500470	Hs.117950	multifunctional polypeptide similar to S	3.01				
55	103036	M13509	Hs.83169	matrix metalloproteinase 1 (Interstitial	27.90				0.70
55	103038	AA926960	Hs.334883 Hs.155324	CDC28 protein kinase 1 matrix metalloproteinase 11 (stromelysin					8.79 4.27
	103060 103099	NM_005940 Al693251	Hs.8248	NADH dehydrogenase (ubiquinone) Fe-S pro		9.80			7.67
	103119	X63629	Hs.2877	cadherin 3, type 1, P-cadherin (placenta	4.05				
C O	103168	X53463	Hs.2704	glutathione peroxidase 2 (gastrointestin	3.07				
60	103185	NM_006825	Hs.74368	transmembrane protein (63kD), endoplasmi		7.40			5.62
	103192 103223	M22440 BE275607	Hs.170009 Hs.1708	transforming growth factor, alpha chaperonin containing TCP1, subunit 3 (g		7.40			4.70
	103242	X76342	Hs.389	alcohol dehydrogenase 7 (class IV), mu o			100.00		
	103316	X83301	Hs.324728	SMA5				9.80	
65	103375	NM_005982	Hs.54416	sine oculis homeobox (Drosophila) homolo	9.71				
	103376	AL036166 NM_007069	Hs.323378 Hs.37189	coated vesicle membrane protein similar to rat HREV107	14.00			11.00	
	103385 103391	X94453	Hs.114366	pyrroline-5-carboxylate synthetase (glut	2.93			11.00	
	103404	BE394784	Hs.78596	proteasome (prosome, macropain) subunit,					5.15
70	103430	BE564090	Hs.20716	translocase of Inner mitochondrial membr				04.40	3.98
	103446 103476	X98834	Hs.79971 Hs.293007	sal (Drosophila)-like 2 aminopeptidase puromycin sensitive		13.00		21.40	
	103470	Y07701 AJ011812	Hs.119018	transcription factor NRF		10.00	6.40		
	103478	BE514982	Hs.38991	S100 calcium-binding protein A2	5.02		•		
75	103515	Y10275	Hs.56407	phosphoserine phosphatase	10.50				
	103558	BE616547	Hs.2785	keratin 17	6.41				3.84
	103580 103587	AA328046 BE270266	Hs.46405 Hs.82128	potymerase (RNA) II (DNA directed) polyp 5T4 oncofetal trophoblast glycoprotein	78.50				J.0 4
	103594	Al368680	Hs.816	SRY (sex determining region Y)-box 2	6.51				
80	103636	NM_006235	Hs.2407	POU domain, class 2, associating factor	3.50				
	103768	AF086009		gb:Homo sapiens full length insert cDNA		0.60			4.48
	103841	AA314821	Hs.38178 Hs.102237	hypothetical protein FLJ23468 tubby super-family protein		8.00 10.40			
_	103847 103913	AF219946 AW967500	Hs.133543	ESTs		. 5.70		15.60	
85	104094	AA418187	Hs.330515	ESTs			6.60		

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	104150	O 02/086 AL122044	44.3 Hs.331633	hypothetical protein DKFZp566N034				26.00	PCT/US02/12476
	104257	BE560621	Hs.9222	estrogen receptor binding site associate		6.80			
	104261	AW248364	Hs.5409	RNA polymerase I subunit					3.98
5	104331	AB040450	Hs.279862	cdk inhibitor p21 binding protein		6.80			
3	104415		Hs.258730	heme-regulated initiation factor 2-alpha	4.21	10.29			
	104558 104590	R56678 AW373062	Hs.88959 Hs.83623	hypothetical protein MGC4816 nuclear receptor subfamily 1, group I, m	4.21			15.79	
	104658	AA360954	Hs.27268	Homo sapiens cDNA: FLJ21933 fis, clone H				17.40	
10	104660	BE298665	Hs.14846	Homo saplens mRNA; cDNA DKFZp564D016 (fr	6.40				
10	104689	AA420450	Hs.292911	ESTs, Highly similar to S60712 band-6-pr					6.55
	104754	AI206234	Hs.155924	cAMP responsive element modulator				10.00	4.42
	104758 104971	BE560269 BE311926	Hs.7010 Hs.15830	NPD002 protein hypothetical protein FLJ12691	2.87				4.47
	105011		Hs.16244	mitotic spindle coiled-coil related prot	3.83				
15	105012		Hs.9329	chromosome 20 open reading frame 1	2.86				
	105026		Hs.124219	hypothetical protein FLJ12934		11.00			
	105076		Hs.37810	hypothetical protein MGC14833					5.01
	105132	AA148164 Al368836	Hs.247280 Hs.24808	HBV associated factor ESTs, Weakly similar to 138022 hypotheti			11.00		3.99
20	105143		Hs.234545	hypothetical protein NUF2R		16.00	11.00		
	105175		Hs.25740	ERO1 (S. cerevisiae)-like	4.32				
	105200	AA328102	Hs.24641	cytoskeleton associated protein 2	3.00				
	105264	AA227934		gb:zr57e08.s1 Soares_NhHMPu_S1 Homo sapi				10.00	
25	105298	BE387790	Hs.26369	hypothetical protein FLJ20287	3.69			0.20	
23	105409 105460	AW505076 AW296078	Hs.301855 Hs.271721	DiGeorge syndrome critical region gene 8 Homo sapiens, clone IMAGE:4179986, mRNA,			7.80	9.20	
	105667		Hs.22030	paired box gene 5 (B-cell lineage specif	4.12		7.00		
	105743	BE246502	Hs.9598	sema domain, Immunoglobulin domain (Ig),	3.82				
20	105782		Hs.57987	B-cell CLL/lymphoma 118 (zinc finger pro			27.00		
30	105848	AW954064	Hs.24951	ESTs			7.60		444
	105891 106019	U55984 AF221993	Hs.289088 Hs.46743	heat shock 90kD protein 1, alpha McKusick-Kaufman syndrome			16.80		4.14
	106069	BE566623	Hs.29899	ESTs, Weakly similar to G02075 transcrip			23.40		
	106073	AL157441	Hs.17834	downstream neighbor of SON	9.50		201.10		
35		AA576953	Hs.22972	hypothetical protein FLJ13352	6.00				
	106159	AK001301	Hs.3487	hypothetical protein FLJ10439					3.95
	106220 106260	D61329 Al097144	Hs.32196 Hs.5250	mitochondrial ribosomal protein L36 ESTs. Weakly similar to ALU1, HUMAN ALU S			13.20		6.04
	106300	Y10043	Hs.19114	high-mobility group (nonhistone chromoso			13.20		5.02
40	106307	AA436174	Hs.37751	ESTs, Weakly similar to putative p150 [6.60			
	106318	AA025610	Hs.9605	cleavage and polyadenylation specific fa					5.04
	106341	AF191020	Hs.5243	hypothetical protein, estradiol-induced			12.00		7.25
	106440 106481	AA449563 D61594	Hs.151393 Hs.17279	glutamate-cysteine ligase, catalytic sub tyrosylprotein sulfotransferase 1	4.75		13.80		
45	106586	AA243837	Hs.57787	ESTs	7.70			10.84	
	106605	AW772298	Hs.21103	Homo saplens mRNA; cDNA DKFZp564B076 (fr				45.60	
	106654	AW075485	Hs.286049	phosphoserine aminotransferase	28,00				
	106785 106813	Y15227 C05766	Hs.20149 Hs.181022	deleted in lymphocytic leukemla, 1 CGI-07 protein	3.00		11.40		
50	106895	AK001826	Hs.25245	hypothetical protein FLJ11269			6.00		•
	106913	Al219346	Hs.86178	M-phase phosphoprotein 9		6.56			
	106919	AW043637	Hs.21766	ESTs, Weakly similar to ALU5_HUMAN ALU S					4.27
	107054		Hs.15978	KIAA1272 protein				34.80	
55	107059 107098	BE614410 Al823593	Hs.23044 Hs.27688	RAD51 (S. cerevisiae) homolog (E coli Re ESTs	4.71			24.80	
55	107104	AU076640	Hs.15243	nucleolar protein 1 (120kD)				24.00	7.05
	107129	AC004770	Hs.4756	flap structure-specific endonuclease 1	2.60		•		
	107198	AV657225	Hs.9846	KIAA1040 protein		19.20			
60	107203	D20426	Hs.41639	programmed cell death 2 DKFZP586E1621 protein	0.50	7.60			
30	107217	AL080235 NM_005629	Hs.35861 Hs.187958	solute carrier family 6 (neurotransmitte	9.50 2.71				
	107218	T74445	Hs.5957	Homo sapiens clone 24416 mRNA sequence			8.71		•
	107516	X57152	Hs.99853	fibrillarin					4.33
65	107529	BE515065	Hs.296585	nucleolar protein (KKE/D repeat)		40.00			4.00 .
03	107728	AA019551	Hs.294151	Homo sapiens, clone IMAGE:3603836, mRNA,		10.80	0.00		
	107851 107901	AA022953 L42612	Hs.61172 Hs.335952	EST keratin 6B	3.40		8.00		
	107922		Hs.61460	Ig superfamily receptor LNIR	2.88				
70	107932	AW392555	Hs.18878	hypothetical protein FLJ21620	7.50				
70	108015	AW298357	Hs.49927	protein kinase NYD-SP15				23.40	
	108056 -108075	AA043675 Al867370	Hs.62633 Hs.139709	ESTs hypothetical protein FLJ12572		•		12.80 12.80	
	108187	BE245374	Hs.27842	hypothetical protein FLJ11210		7.00		12.00	
	108296	N31256	Hs.161623	ESTs		6.60			
75	108305	AA071391		gb:zm61e06.r1 Stratagene fibroblast (937				11.80	
	108393	AA075211	Un COACE	gb:zm86a08.r1 Stratagene ovarian cancer				11.80	
	108480 108554	AL133092 AA084948	Hs.68055	hypothetical protein DKFZp434I0428 gb:zn13b09.s1 Stratagene hNT neuron (937		6.40		20.80	
	108573	AA086005		gb:zi84c04.s1 Stratagene colon (937204)		J.73		25.40	
80	108584	AA088326	Hs.120905	Homo saplens cDNA FLJ11448 fis, clone HE		9.60			
	108597	AK000292	Hs.278732	hypothetical protein FLJ20285	0.00			14.60	
	108695	AB029000	Hs.70823 Hs.70832	KIAA1077 protein ESTs	3.00			10.00	
	108699 108700	AA121514 AA121518	Hs.193540	ESTs, Moderately similar to 2109260A B c			11.00	10.00	
85	108780	AU076442	Hs.117938	collagen, type XVII, alpha 1	11.21				
				· · · · · ·					

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	108810	AW295647	Hs.71331	hypothetical protein MGC5350	8.50	- 40			
	108816 108857	AA130884 AK001468	Hs.270501 Hs.62180	ESTs, Moderately similar to ALU2_HUMAN anillin (Drosophila Scraps homolog), act	4.00	7.40			
,	108860	AA133334	Hs.129911	ESTs	6.09				
5	108937	AL050107	Hs.24341	transcriptional co-activator with PDZ-bi	3.00				
	109010	NM_007240	Hs.44229	dual specificity phosphatase 12	2.69		•		4.50
	109121 109166	BE389387 AA219691	Hs.49767 Hs.73625	NADH dehydrogenase (ubiquinone) Fe-S pro RAB6 interacting, kinesin-like (rabkines	10.58				4.53
	109100	AA766998	Hs.85874	Human DNA sequence from clone RP11-16L21	10.55	9.00			
10	109415	U80736	Hs.110826	trinucleotide repeat containing 9		51.40			
	109418	AI866946	Hs.161707	ESTs			47.00	11.00	
	109454	AA232255 AW967069	Hs.295232 Hs.211556	ESTs, Moderately similar to A46010 X-II hypothetical protein MGC5487			17.60 9.49		
	109502 109543	AA564994	Hs.222851	ESTs .		12.67	5.45		
15	109648	H17800	Hs.7154	ESTs				10.40	
	109680	AB037734	Hs.4993	KIAA1313 protein			33.20	40.00	
	109700 109704	F09609 Al743880	Hs.12876	gb:HSC33H092 normalized infant brain cDN ESTs			11.00	16.00	
	109792	R49625	113.12070	gb:yg61f03.s1 Soares infant brain 1NiB H				12.60	
20	109981	BE546208	Hs.26090	hypothetical protein FLJ20272	4.00				
	109998	AL042201	Hs.21273	transcription factor NYD-sp10		7.80			
	110039 110156	H11938 AA581322	Hs.21907 Hs.4213	histone acetyltransferase hypothetical protein MGC16207		7.00			4.24
	110500	AA907723	Hs.36962	ESTs	4.50				
25	110551	AW450381	Hs.14529	ESTs	0.00	8.60			
	110561 110854	AA379597 BE612992	Hs.5199 Hs.27931	HSPC150 protein similar to ubiquitin-con hypothetical protein FLJ10607 similar to	3.06	6.80			
	110886	AW274992	Hs.72249	three-PDZ containing protein similar to		0.00	8.80		
20	110916	BE178102	Hs.24349	ESTs		6.80		40.00	
30	111003	N52980	Hs.83765 Hs.263925	dihydrofolate reductase US1-interacting protein NUDE1, rat homo	2.54			16.80	
	111337 111434	AA837396 R01608	Hs.142736	ESTs	2.04			9.80	
	111439	A1476429	Hs.19238	ESTs				10.40	
25	111540	U82670	Hs.9786	zinc finger protein 275			15.40	0.20	
35	111597 111895	R11499 T80581	Hs.189716 Hs.12723	ESTs Homo sapiens clone 25153 mRNA sequence		6.80		9.20	
	111929	AF027208	Hs.112360	prominin (mouse)-like 1				14.67	
	112054	R43590		gb:yc85g02.s1 Soares Infant brain 1NiB H		10.80		40.00	
40	112210 112244	R49645 AB029000	Hs.7004 Hs.70823	ESTs KIAA1077 protein	2.99			10.20	
	112382	R59904	110.7 0020	gb:yh07g12.s1 Soares infant brain 1NIB H		6.60			
	112392	R60763	Hs.193274	ESTs, Moderately similar to 157588 HSrel			7.10		
	112442 112539	AA280174 R70318	Hs.285681 Hs.339730	Williams-Beuren syndrome chromosome regi ESTs	3.00			37.20	
45	112772	A1992283	Hs.35437	ESTs, Moderately similar to 138026 MLN 6				14.60	
	112869	BE261750	Hs.4747	dyskeratosis congenita 1, dyskerin					4.83
	112935 112970	R71449 AA694010	Hs.268760 Hs.6932	ESTs Homo sapiens clone 23809 mRNA sequence	2.73			12.00	
	112973	AB033023	Hs.318127	hypothetical protein FLJ10201	11.50			12.00	
50	112992	AL157425	Hs.133315	Homo saplens mRNA; cDNA DKFZp761J1324 (f	45.00		10.89		
	113063 113073	W15573 N39342	Hs.5027 Hs.103042	ESTs, Weakly similar to A47582 B-cell gr microtubule-associated protein 18	15.00		15.31		
	113078	T40444	Hs.118354	CAT56 protein		7.00			·
55	113238	R45467	Hs.189813	ESTs				41.20	
55	113591 113702	T91881 T97307	Hs.200597	KIAA0563 gene product qb:ye53h05,s1 Soares fetal liver spleen	25.00			9.40	
	113844	Al369275	Hs.243010	Homo sapiens cDNA FLJ14445 fis, clone HE	25.00			13.91	
	113984	R96696	Hs.35598	ESTs		7.80			
60	114073 114162	R44953 AF155661	Hs.22908 Hs.22265	Homo sapiens mRNA; cDNA DKFZp434J1027 (f pyruvate dehydrogenase phosphatase	3.42	7.20			
00	114208	AL049466	Hs.7859	ESTs	0.42		6.74		
	114251	H15261	Hs.21948	ESTs				33.20	
	114285	R44338	Hs.22974	ESTs ESTs				13.20 10.00	
65	114313 114339	H18456 AA782845	Hs.27946 Hs.22790	ESTS		7.80		10.00	
-	114407	BE539976	Hs.103305	Homo sapiens mRNA; cDNA DKFZp434B0425 (f					4.14
	114560	AI452469	Hs.165221	eSTs ab:zn90d09.r1 Stratagene lung carcinoma		7.60		9.80	
	114699 114767	AA127386 Al859865	Hs.154443	minichromosome maintenance deficient (S	3.21	7.00			
70	114793	AA158245		gb:zo76c03.s1 Stratagene pancreas (93720			6.00		
	114833	Al417215	Hs.87159	hypothetical protein FLJ12577 chaperonin containing TCP1, subunit 6A (11.40	4.31
	115047 115060	BE270930 AF052693	Hs.82916 Hs.198249	gap junction protein, beta 5 (connexin 3					4.03
	115097	AA256213	Hs.72010	ESTs				35.40	
75	115113	AA256460		gb:zr81a04.s1 Soares_NhHMPu_S1 Homo sapi				15.20	4.40
	115123 115134	AA256641 AW968073	Hs.236894 Hs.194331	ESTs, Highly similar to S02392 alpha-2-m ESTs, Highly similar to A55713 inositol				12.40	4.19
	115291	BE545072	Hs.122579	hypothetical protein FLJ10461	25.00				
00	115347	AA356792	Hs.334824	hypothetical protein FLJ14825		7.00			
80	115414	AA662240	Hs.283099	AF15q14 protein	3,25 3.68				
	115522 115536	BE614387 AK001468	Hs.333893 Hs.62180	c-Myc target JPO1 anillin (Drosophila Scraps homolog), act	3.66 10.50				
	115566	AJ142336	Hs.43977	Human DNA sequence from clone RP11-196N1				24.40	
85	115645 115648	Al207410 AW016811	Hs.69280 Hs.234478	Homo sapiens, clone IMAGE:3636299, mRNA, Homo sapiens cDNA: FLJ22648 fis, clone H	4.17		6.00		
0.5	113040	V14010011	· 101.20***10	From achiena actar i estrada noi arano (1			5.44		

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	115652	BE093589	Hs.38178	hypothetical protein FLJ23468	3.81	•			
	115697	D31382	Hs.63325	transmembrane protease, serine 4	62.14			44.00	
	115793	AA424883	Hs.70333	hypothetical protein MGC10753				11.80 9.71	
5	115816 115892	BE042915 AA291377	Hs.287588 Hs.50831	Homo sapiens cDNA FLJ13675 fis, clone PL ESTs			27.40	5.7 1	
J .	115906	Al767756	Hs.82302	Homo sapiens cDNA FLJ14814 fis, clone NT	2.53				
	115909	AW872527	Hs.59761	ESTs, Weakly similar to DAP1_HUMAN DEATH	11.82			24.20	
	115965	AA001732	Hs.173233	hypothetical protein FLJ10970				34.29	8.23
10	115978 115985	AL035864 AA447709	Hs.69517 Hs.268115	cDNA for differentially expressed CO16 g ESTs, Wealdy similar to T08599 probable	3.00				0.23
10	116090	AI591147	Hs.61232	ESTs	5.17				
	116096	AA682382	Hs.59982	ESTs		40.00	8.20		
	116127	AF126743	Hs.279884	DNAJ domain-containing		10.60			5.82
15	116157 116190	BE439838 A1949095	Hs.44298 Hs.67776	mitochondrial ribosomal protein S17 ESTs, Weakly similar to T22341 hypotheti					4.08
13	116278	NM_003686	Hs.47504	exonuclease 1	9.50				
	116335	AK001100	Hs.41690	desmocollin 3	3.67	7.00			
	116496	AW450694	Hs.21433	hypothetical protein DKFZp547J036		7.00		12.60	
20	116503 116674	AI925316 AI768015	Hs.212617 Hs.92127	ESTs ESTs			32.00	12.00	
20	116929	AA586922	Hs.80475	polymerase (RNA) II (DNA directed) polyp		7.60			
	116973	AJ702054	Hs.166982	phosphatidylinositol glycan, class F		9.80		10.20	
	116993 117079	Al417023 H92325	Hs.40478	ESTs gb:ys85f05.s1 Soares retina N2b4HR Homo				15.20	
25	117317	Al263517	Hs.43322	ESTs				13.40	
	117326	N23629	Hs.241420	Homo sapiens mRNA for KIAA1756 protein,				20.60	
	117396	W20128	Hs.296039	ESTs				10.60 16.00	
	117412	N32536 N32528	Hs.42645 Hs.146286	ESTs kinesin family member 13A				9.11	
30	117519 117693	AW179019	Hs.112110	mitochondrial ribosomal protein L42				••••	4.01
	117721	N46100	Hs.93939	EST				19.80	
	117881	AF161470	Hs.260622	butyrate-induced transcript 1	2.71			17.80	
	117903	AA768283	Hs.47111 Hs.172089	ESTs Homo sapiens mRNA; cDNA DKFZp586l2022 (f				17.00	4,17
35	117992 118013	Al015709 Al674126	Hs.94031	ESTs				10.60	
50	118017	AI813444	Hs.42197	ESTs			8.82		
	118186	N22886	Hs.42380	ESTs		7.00		13.80	
	118325 118367	A1868065 N64269	Hs.166184 Hs.48946	intersectin 2 EST			6.14	15.00	
40	118368	N64339	Hs.48956	gap junction protein, beta 6 (connexin 3	3.14				
	118472	AL157545	Hs.42179	bromodomain and PHD finger containing, 3			12.40	40.00	
	118709	AA232970	Hs.293774 Hs.55209	ESTs Homo sapiens mRNA; cDNA DKFZp434K0514 (f	4.50			12.20	•
	119025 119027	BE003760 AF086161	Hs.114611	hypothetical protein FLJ11808	3.22				
45	119052	R10889	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	gb:yf38d02.s1 Soares fetal liver spleen		9.60			
	119164	AF221993	Hs.46743	McKusick-Kaufman syndrome			6.60	10.80	
	119186 119243	Al979147 T12603	Hs.101265	hypothetical protein FLJ22593 gb:CHR90123 Chromosome 9 exon il Homo sa				9.44	
	119490	AA195276	Hs.263858	ESTs, Moderately similar to B34087 hypot				11.80	
50	119499	Al918906	Hs.55080	ESTs		40.00	14.80		
	119599	W45552	Un 404204	gb:zc26d03.s1 Soares_senescent_fibroblas hypothetical protein	17.00	12.60			
	119780 119845	NM_016625 W79123	Hs.191381 Hs.58561	G protein-coupled receptor 87	13.50				
	119941	AA699485	Hs.58896	ESTs		8.00			
55	119994	AA642402	Hs.59142	ESTs	7.73		39.60		•
	120102 120104	W67353 AK000123	Hs.170218 Hs.180479	KIAA0251 protein hypothetical protein FLJ20116	2.91		39.60		•
	120294	AK000059	Hs.153881	Homo saplens NY-REN-62 antigen mRNA, par			8.20		
C 0	120486	AW368377	Hs.137569	tumor protein 63 kDa with strong homolog	8.73				
60	120599	AA804448	Hs.104463 Hs.97258	ESTs ESTs, Moderately similar to S29539 ribos		7.00		10.00	
	120699 120715	A1683243 AA292700	NS.97 200	obizs59a06 st NCL CGAP GCR1 Homo saplens		9.40		10.00	
	120821	Y19062	Hs.96870	staufen (Drosophila, RNA-binding protein				13.80	
65	120859	AA826434	Hs.1619	achaete-scute complex (Drosophila) homol		9.00			
65	120880	AA360240	Hs.97019 Hs.97587	EST EST		15.60	27.66		
	120983 121034	AA398209 AL389951	Hs.271623	nucleoporin 50kD			20.80		
	121121	AA399371	Hs.189095	similar to SALL1 (sal (Drosophila)-like		22,80			
70	121313	AA402713	Hs.97872	ESTs	25.71			10.00	
70	121369 121376	AW450737 AA448103	Hs.128791 Hs.187958	CGI-09 protein solute carrier family 6 (neurotransmitte	20.71				5.42
	121476	AA412311	Hs.97903	ESTs		8.30			
	121509	AA868939	Hs.97888	ESTs	40.50	8.59			
75	121553 121753	AA412488 AK000552	Hs.48820 Hs.323518	TATA box binding protein (TBP)-associat WD repeat domain 5	18.50 7.00				
75	121838	AA425680	Hs.98441	ESTS				10.40	
	121857	BE387162	Hs.280858	ESTs, Highly similar to A35661 DNA excis	6.00			40.00	
	121991	AA430058	Hs.98649	EST			8.60	12.20	
80	122089 122105	AW016543 AW241685	Hs.98682 Hs.98699	hypothetical protein FKSG32 ESTs			6.14		
5 0	122163	AA435702	Hs.98829	EST				10.40	
	122318	AA429743		gb:zv60b05.r1 Soares_testis_NHT Homo sap	10.50			18.20	
	122335 122338	AA443258 AA443311	Hs.241551 Hs.98998	chloride channel, calcium activated, fam ESTs	13.50 4.80				
85	122336		Hs.99087	ESTs, Weakly similar to S47073 finger pr		8.00			
				•					

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	122512	AF053305	Hs.98658	budding uninhibited by benzimidazoles 1			8.80	9.40	
	122516. 122702	AA449352 Al220089	Hs.99217 Hs.99439	ESTs ESTs		9.20		9.40	
_	122852	A1580056	Hs.98992	ESTs				10.40	
5	122925	AW268962	Hs.111335	ESTs		6.80	12.60		
	123005 123044	AW369771 AK001035	Hs.52620 Hs.130881	integrin, beta 8 B-cell CLL/lymphoma 11A (zinc finger pro			12.00		5.35
	123160	AA488687	Hs.284235	ESTs, Weakly similar to 138022 hypotheti			6.06		
10	123315	AA496369		gb:zv37d10.s1 Soares overy tumor NbHOT H			12.40 11.80		
10	123329 123497	Z47542 AA765256	Hs.179312 Hs.135191	small nuclear RNA activating complex, po ESTs, Weakly similar to unnamed protein		12.00	11.00		
•	123518	AL035414	Hs.21068	hypothetical protein			13.00		
	123519	AW015887	Hs.112574	ESTs		12.20	7.80		
15	123614 123616	AK000492 AA680003	Hs.98806 Hs.109363	hypothetical protein Homo sapiens cDNA: FLJ23603 fis, clone L			7.00	10.60	
13	123673	BE550112	Hs.158549	ESTs, Weakly similar to T2D3_HUMAN TRANS	23.00				
	123727	AI083986	Hs.282977	hypothetical protein FLJ13490		7.00	9.80		
	123731 123752	AA609839 AA227714	Hs.179703	gb:ae62f01.s1 Stratagene lung carcinoma KIAA0129 gene product	3.50		5.00		
20	123900	AA621223	Hs.112953	EST				12.80	
	124006	Al147155	Hs.270016	ESTs	97.00				
	124059 124069	BE387335 AF134160	Hs.283713 Hs.7327	ESTs, Weakly similar to S64054 hypotheti claudin 1	3.02		27.80		
	124191	T96509	Hs.248549	ESTs, Moderately similar to S65657 alpha				35.80	
25	124273	AA457211	Hs.8858	bromodomain adjacent to zinc finger doma		7.20		11.00	
	124297 124305	AL080215 AW963221	Hs.102301	Homo sapiens mRNA; cDNA DKFZp586J0323 (f gb:EST375294 MAGE resequences, MAGH Homo				16.00	
	124676	Al360119.com	pHs.181013	phosphoglycerate mutase 1 (brain)					6.08
20	124874	BE550182	Hs.127826	RaiGEF-like protein 3, mouse homolog		9.40		21.00	
30	124904 124969	AK000483 AI650360	Hs.93872 Hs.100256	KIAA1682 protein ESTs		3.40		10:80	
	125000	T58615	Hs.110640	ESTs				9.80	
	125201	AA693960	Hs.103158	ESTs, Weakly similar to T33296 hypotheti		7.60 6.59			
35	125266 125299	W90022 T32982	Hs.186809 Hs.102720	ESTs, Highly similar to LCT2_HUMAN LEUKO ESTs		0.00		9.57	
-	125356	AI057052	Hs.133554	ESTs, Weakly similar to Z195_HUMAN ZINC				14.00	
	125370	AA256743	Hs.134158	Homo sapiens, Similar to KIAA0092 gene p			8.20	13.20	•
	125418 125433	AA777690 AL162066	Hs.188501 Hs.54320	ESTs hypothetical protein DKFZp762D096		21.40		10.50	
40	125437	A1609449	Hs.140197	ESTs		6.96			
	125446	BE219987	Hs.166982	phosphatidylinositol glycan, class F hypothetical protein AF140225		8.80		11.20	
	125711 125756	AA305800 BE174587	Hs.5672 Hs.289721	growth arrest specific transcript 5					4,31
4.5	125757	Al274906	Hs.166835	ESTs, Highly similar to 1814460A p53-ass	0.00			15.60	
45	125769 125839	BE270266 AW836261	Hs.82128 Hs.337717	5T4 oncofetal trophoblast glycoprotein ESTs	3.20	8.20			
	125850	W85858	Hs.99804	ESTs	2.65				•
	125875	H14480		gb:ym18b09.r1 Soares infant brain 1NIB H		7.40			4.23
50	125924 125972	BE272506 Al927475	Hs.82109 Hs.35406	syndecan 1 ESTs, Highly similar to unnamed protein					3.98
50	126034	H60340	113.00400	gb:yr39b04.r1 Soares fetal liver spleen				10.60	
	126327	AA432266	Hs.44648	ESTs		11.60 6.67			
	126345 126435	N49713 AW614529	Hs.285847	gb:yv23f06.s1 Soares fetal liver spleen CGI-19 protein		0.01		10.60	
55	126487	AA283809	Hs.184601	solute carrier family 7 (cationic amino					4.38
	126521	AI475110	Hs.203933	ESTs gb:zc76d03.s1 Pancreatic Islet Homo sapl		6.60		14.80	
	126522 126543	W31912 AL035864	Hs.69517	cDNA for differentially expressed CO16 g					4.01
CO	126567	AA058394	Hs.57887	ESTs, Weakly similar to KIAA0758 protein			7.80	11.60	
60	126605 126627	AA676910 AA497044	Hs.20887	gb:zj65h07.s1 Soares_fetat_liver_spleen_ hypothetical protein FLJ10392				14.60	
	126628	N49776	Hs.170994	hypothetical protein MGC10946	8.00				
	126737	AW976516	Hs.283707	Homo sapiens cDNA: FLJ21354 fis, clone C	2.92 7.50				
65	126795 126802	AW975076 AW805510	Hs.172589 Hs.97056	nuclear phosphoprotein similar to S. cer hypothetical protein FLJ21634	7.50	11.60			
Ų,	126892	AF121856	Hs.284291	sorting nexin 6	3.50				
	126928	AA480902	Hs.137401	ESTs gb:zq89h10.r1 Stratagene hNT neuron (937				22.83 11.80	
	126979 126986	AA210954 Al279892	Hs.46801	sorting nextn 14				11.60	
70	126992	Al809521	•	gb:wf30e03.x1 Soares_NFL_T_GBC_S1 Homo s				20.80	
	127066	R25066		gb:yg42c07.r1 Soares Infant brain 1NIB H gb:EST54026 Fetal heart II Homo saplens				27.60 21.60	
	127099 127139	AA347668 AA830233	Hs.293585	ESTs				11.20	
75	127209	AA305023	Hs.81964	SEC24 (S. cerevislae) related gene famil	3.10				
75	127221 127225	BE062109 AA315933	Hs.241551 Hs.120879	chloride channel, calcium activated, fam ESTs	2.76			16.80	
	127313	AK002014	Hs.47546	Homo sapiens cDNA FLJ11458 fis, clone HE	14.00				•
	127444	AW978474	Hs.7560	Homo sapiens mRNA for KIAA1729 protein,		11.20		13.60	
80	127500 127524	AW971353 Al243596	Hs.162115 Hs.94830	ESTs ESTs, Moderately similar to T03094 A-kin			7.80		
	127540	N45572	Hs.105362	Homo saplens, clone MGC:18257, mRNA, com	3.53			12.00	
	127599	AA613204 X80031	Hs.150399 Hs.530	ESTs collagen, type IV, alpha 3 (Goodpasture	•			13.80 28.00	
	127609 127662	W80755	Hs.8294	KIAA0196 gene product				19.80	
85	127668	Al343257	Hs.139993	ESTs				11.20	

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	127746		Hs.120189	ESTs				14.18	
		AA741368	Hs.291434	ESTs	4.50			24.60	
	127817	AA836841 Al302471	Hs.163085 Hs.124292	ESTs Homo sapiens cDNA: FLJ23123 fis, clone L				24.60 9.20	•
5	127959 127960	Al613226	Hs.41569	phosphatidic acid phosphatase type 2A				16.83	
_	127969	F06498	Hs.93748	Homo sapiens cDNA FLJ14676 fis, clone NT		13.60			
	128015	Z21169	Hs.334659	hypothetical protein MGC14139		7.00		37.40	
	128027 128077	Al433721 Al310330	Hs.164153 Hs.128720	ESTs ESTs				9.60	
10	128168	NM_006147	Hs.11801	Interferon regulatory factor 6				9.24	
	128226	Al284940	Hs.289082	GM2 ganglioside activator protein	19.00				
	128305	A1954968	Hs.279009	matrix Gla protein		9.00		10.40	•
	128341 128527	AA191420 AA504583	Hs.185030 Hs.101047	ESTs transcription factor 3 (E2A immunoglobul		5.00			4.30
15	128539	R46163	Hs.258618	ESTs		12.60			
	128568	H12912	Hs.274691	adenylate kinase 3				40.00	4.56
	128572		Hs.256583	interleukin enhancer binding factor 3, 9 cysteine and glycine-rich protein 2			16.80	10.00	
	128777 128781	AI878918 N71826	Hs.10526 Hs.105465	small nuclear ribonucleoprotein polypept			10.00		4.48
20	128796		Hs.105924	defensin, beta 2		8.12			
	128920	AA622037	Hs.166468	programmed cell death 5					4.62 4.04
	128924 128971	BE279383 H05132	Hs.26557 Hs.107510	plakophilin 3 ESTs		12.60			4.04
	129008	AL079648	Hs.301088	ESTs		8.80			
25	129041	BE382756	Hs.169902	solute carrier family 2 (facilitated glu	0.50				6.05
	129075 129105	BE250162	Hs.83765 Hs.108681	dihydrofolate reductase Homo sapiens brain tumor associated prot	2.59		6.67		
	129189	Al769160 AB023179	Hs.9059	KIAA0962 protein		8.00	0.01		
	129229	AF013758	Hs.109643	polyadenylate binding protein-interactin	4.00	•			
30	129241	A1878857	Hs.109706	hematological and neurological expressed	0 55				4.06
	129300 129404	W94197 Al267700	Hs.110165 Hs.317584	ribosomal protein L26 homolog ESTs	2.55 18.00				
	129457	X61959	Hs.207776	aspartylglucosaminidase	6.50				
25	129466	L42583	Hs.334309	keratin 6A	12.94			44.00	
35	129494 129605	AI148976	Hs.112062 Hs.115947	ESTs keratin 16 (focal non-epidermolytic palm				11.00	4.46
	129641	AF061812 Al911527	Hs.11805	ESTs				12.00	
	129665	AW163331	Hs.118778	KDEL (Lys-Asp-Glu-Leu) endoplasmic retic		•			4.70
40	129703	BE388665	Hs.179999	Homo sapiens, clone IMAGE:3457003, mRNA					4.02 5.71
40	129720 129748	AA156214 M16707	Hs.12152 Hs.123053	APMCF1 protein H4 histone, family 2	3.50				0.71
	129890	AI868872	Hs.282804	hypothetical protein FLJ22704					4.21
	129896	BE295568	Hs.13225	UDP-Gal:betaGicNAc beta 1,4- galactosylt	2.56				4.03
45	129945 130010	BE514376 AA301116	Hs.165998 Hs.142838	PAI-1 mRNA-binding protein nucleolar phosphoprotein Nopp34			7.00		4.03
7.5	130026	T40480	Hs.332112	EST		6.40			
	130080	X14850	Hs.147097	H2A histone family, member X	2.74				4.65
	130149 130285	AW067805 AA063546	Hs.172665 Hs.75981	methylenetetrahydrofolate dehydrogenase ubiquilin specific protease 14 (tRNA-gua	2.14		7.40		
50	130441	U63630	Hs.155637	protein kinase, DNA-activated, catalytic					3.91
	130482	AW409701	Hs.1578	baculoviral IAP repeat-containing 5 (sur	4.87			9.60	
	130500 130524	AB007913 U89995	Hs.158291 Hs.159234	KIAA0444 protein forkhead box E1 (thyroid transcription f			13.40	9.00	
	130541	X05608	Hs.211584	neurofilament, light polypeptide (68kD)			8.20		
55	130553	AF062649	Hs.252587	pituitary tumor-transforming 1			7.00		6.06
	130567 130577	AA383092 M69241	Hs.1608 Hs.162	replication protein A3 (14kD) Insulin-like growth factor binding prote	3.04		7.00		
	130627	BE003054	Hs.1695	matrix metalloproteinase 12 (macrophage	3.87				
~	130648	Al458165	Hs.17296	hypothetical protein MGC2376				16.20	
60	130697	L29472	Hs.1802 Hs.18747	major histocompatibility complex, class POP7 (processing of precursor, S. cerevi				17.80	5.28
	130744 130800	H59696 Al187292	Hs.19574	hypothetical protein MGC5469					4.43
	130867	NM_001072	Hs.284239	UDP głycosyltransferase 1 family, polype	16.84				4.00
65	130869	J03626	Hs.2057	uridine monophosphate synthetase (orotat				9.60	4.92
03	130925 130994	ÀF093419 W17044	Hs.169378 Hs.327337	multiple PDZ domain protein ESTs		12.40		5.00	
	131028	Al879165	Hs.2227	CCAAT/enhancer binding protein (C/EBP),	10.21				
	131031	NM_001650	Hs.288650	aquaporin 4				9.80 9.60	
70	131041 131058	T15767 W28545	Hs.22452 Hs.101514	Homo sapiens mRNA for KIAA1737 protein, hypothetical protein FLJ10342				17.00	
, 0	131090	Al143139	Hs.2288	visinin-like 1	2.74				
	131112	H15302	Hs.168950	Homo saplens mRNA; cDNA DKFZp566A1046 (f	0.40		8.80		
	131148 131185	AW953575 BE280074	Hs.303125 Hs.23960	p53-induced protein PIGPC1 cyclin B1	3.12 3.07				
75	131200	BE540516	Hs.293732	hypothetical protein MGC3195	3.07				
-	131219	W25005	Hs.24395	small inducible cytokine subfamily B (Cy	2.87			44.00	
	131257	AW339037	Hs.24908 Hs.143134	ESTs ESTs			19.20	14.67	
	131375 131460	AW293165 NM_003729	Hs.27076	RNA 3'-terminal phosphate cyclase	3.50				
80	131476	AI521663	Hs.334644	hypothetical protein FLJ14668	15.00				
	131510	BE245374	Hs.27842	hypothetical protein FLJ11210 MRS2 (S. cerevisiae)-like, magnesium hom			7.80 7.00		
	131646 131786	BE302464 BE000971	Hs.30057 Hs.306083	Novel human gene mapping to chomosome 22	2.65		7.00		
0.5	131839	AB014533	Hs.33010	KIAA0633 protein	-			35.20	4.44
85	131843	AA192315	Hs.184062	putative Rab5-interacting protein					4.11

	W	O 02/086	443						PCT/US02/12476
	131877	J04088	Hs.156346	topoisomerase (DNA) II alpha (170kD)	19.00				
	131885	BE502341	Hs.3402	ESTs	6.48		8.40		
	131921 131945	AA456093 NM_002916	Hs.34720 Hs.35120	ESTs replication factor C (activator 1) 4 (37	56.00		0.40		
5	131958	NM_014062	Hs.3566	ART-4 protein					3.82
	131965	W79283	Hs.35962	ESTs	3.03				
	132000 132040	AW247017 NM_001196	Hs.36978 Hs.315689	melanoma antigen, family A, 3 Homo sapiens cDNA: FLJ22373 fls, clone H	3.30	9.80			
	132109	AW190902	Hs.40098	cysteine knot superfamily 1, BMP antagon	21.00				
10	132114	NM_006152	Hs.40202	lymphoid-restricted membrane protein		8.40	•		
	132162	AA315805	Hs.94560	desmoglein 2	2.70				12.25
	132164 132180	AI752235 NM_004460	Hs.41270 Hs.418	procollagen-lysine, 2-oxoglutarate 5-dlo fibroblast activation protein, alpha	2.70 2.71				
	132181	AW961231	Hs.16773	Homo saplens clone TCCCIA00427 mRNA sequ	3.83				
15	132182	NM_014210	Hs.70499	ecotropic viral Integration site 2A				13.20	
	132231 132277	AA662910 AK001745	Hs.42635 Hs.184628	hypothetical protein DKFZp434K2435 hypothetical protein FLJ10883	9.50 4.50				
	132328	NM_014787	Hs.44896	DnaJ (Hsp40) homolog, subfamily B, membe	4.00			9.20	
••	132394	AK001680	Hs.30488	DKFZP434F091 protein				19.80	
20	132424	AA417878	Hs.48401	ESTs, Moderately similar to ALU8_HUMAN A			8.60		•
	132528 132543	T78736 BE568452	Hs.50758 Hs.5101	SMC4 (structural maintenance of chromoso protein regulator of cytokinesis 1	4.38		27.40		
	132544	L19778	Hs.51011	H2A histone family, member P		7.00			
25	132550	AW969253	Hs.170195	bone morphogenetic protein 7 (osteogenic	2.64			45.00	
25	132552 132581	BE621985 AK000631	Hs.296922 Hs.52256	thiopurine S-methyltransferase hypothetical protein FLJ20624			6.60	15.83	
	132617	AF037335	Hs.5338	carbonic anhydrase XII	4.95		0.00		
	132638	A1798870	Hs.54277	DNA segment on chromosome X (unique) 992		8.20			
30	132653	Z15008	Hs.54451	laminin, gamma 2 (nicein (100kD), kalini	4.38				4.36
30	132669 132710	W38586 W74001	Hs.293981 Hs.55279	guanine nucleotide binding protein (G pr serine (or cysteine) proteinase inhibito	4.60				4.00
	132771	Y10275	Hs.56407	phosphoserine phosphatase	3.71				
	132799	W73311	Hs.169407	SAC2 (suppressor of actin mutations 2,				9.48	5.83
35	132833 132892	U78525 AW834050	Hs.57783 Hs.9973	eukaryotic translation initiation factor tensin				12.00	0.00
	132906	BE613337	Hs.234896	geminin	3.09				
	132959	AW014195	Hs.61472	ESTs, Weakly similar to YAE6_YEAST HYPOT	3.50				3.87
	132962 132990	AA576635 X77343	Hs.6153 Hs.334334	CGI-48 protein transcription factor AP-2 alpha (activat	6.18				
40	132994	AA112748	Hs.279905	clone HQ0310 PRO0310p1	3.19				
	133000	AL042444	Hs.62402	p21/Cdc42/Rac1-activated kinase 1 (yeast	2.96 2.55				
	133050 133083	X73424 BE244588	Hs.63788 Hs.6456	propionyl Coenzyme A carboxylase, beta p chaperonin containing TCP1, subunit 2 (b	2.55				4.00
4 =	133086	L17131	Hs.139800	high-mobility group (nonhistone chromoso					8.96
45	133134	AF198620	Hs.65648	RNA binding motif protein 8A				10.80	4.28
	133155 133181	M58583 X91662	Hs.662 Hs.66744	cerebellin 1 precursor twist (Drosophila) homolog (acrocephalos	3.00			10.00	
	133204	BE267696	Hs.254105	enolase 1, (alpha)					4.63
50	133412 133421	U41493 AF134160	Hs.73112 Hs.7327	guanine nucleotide binding protein (G pr claudin 1	2.85	12.50			
30	133451	AW970026	Hs.73818	ublquinol-cytochrome c reductase hinge p	2.00				4.66
	133453	A1659306	Hs.73826	protein tyrosine phosphatase, non-recept		6.80			
	133504 133506	NM_004415 BE562958	Hs.74316 Hs.74346	desmoplakin (DPI, DPII) hypothetical protein MGC14353	6.14				4.55
55	133615	M62843	Hs.75236	ELAV (embryonic lethal, abnormal vision,				17.80	
	133627	NM_002047	Hs.75280	glycyl-tRNA synthetase					4.85
	133649 133669	U25849 NM_006925	Hs.75393 Hs.166975	acid phosphatase 1, soluble splicing factor, arginine/serine-rich 5				14.00	6.34
	133749	L20852	Hs.100370	solute carrier family 20 (phosphate tran			6.11		
60	133776	BE268649	Hs.177766	ADP-ribosyltransferase (NAD+; poly (ADP-	0.07				4.91
	133865 133946	AB011155 AJ001258	Hs.170290 Hs.173878	discs, large (Drosophila) homolog 5 NIPSNAP, C. elegans, homolog 1	3.07				4.60
	133973	N55540	Hs.78026	ESTs, Weakly similar to similar to ankyr				13.00	
CE	134047	BE262529	Hs.78771	phosphoglycerate kinase 1					3.85
65	134098 134107	BE513171 NM_005629	Hs.79086 Hs.187958	mitochondrial ribosomal protein L3 solute carrier family 6 (neurotransmitte	2.56		8.20		
	134112	AW449809	Hs.79150	chaperonin containing TCP1, subunit 4 (d			0.20		4.08
	134158	U15174	Hs.79428	BCL2/adenovirus E1B 19kD-Interacting pro	31.00		04.00		
70	134160 134168	T98152 AA398908	Hs.79432 Hs.181634	fibrillin 2 (congenital contractural ara Homo sapiens cDNA: FLJ23602 fis, clone L			24.60		6.71
, 0	134185	AA285136	Hs.301914	neuronal specific transcription factor D				14.74	
	134201	L35035	Hs.79886	ribose 5-phosphate isomerase A (ribose 5	4 50	8.40			
	134272 134276	X76040 BE083936	Hs.278614 Hs.80976	protease, serine, 15 antigen identified by monoclonal antibod	4.50	9.00			
75	134353	AL138201	Hs.82120	nuclear receptor subfamily 4, group A, m				16.40	
	134367	AA339449	Hs.82285	phosphoribosylglycinamide formyltransfer	2.80 4.68				
	134380 134423	AU077143 H53497	Hs.179565 Hs.83006	minichromosome maintenance deficient (S. CGI-139 protein	4.00				3.84
00	134469	AA279661	Hs.83753	small nuclear ribonucleoprotein polypept					5.81
80	134470	X54942	Hs.83758	CDC28 protein kinase 2					4.21 7.30
	134498 134502	AW246273 BE148534	Hs.84131 Hs.84168	threonyl-tRNA synthetase UV-B repressed sequence, HUR 7		13.60		•	
	134510	NM_002757	Hs.250870	mitogen-activated protein kinase kinase				9.70	4.69
85	134548 134654	N95406 AK001741	Hs.333495 Hs.8739	Deleted in split-hand/split-foot 1 regio hypothetical protein FLJ10879	6.00		•		4.63
55	107004		10.0100						

	W	O 02/086	443					PCT/US02/12476		
	134724	AF045239	Hs.321576	ring finger protein 22				12.00		
	134743	AA044163	Hs.89463	potassium large conductance calcium-acti	4.00					
	134781	AA374372	Hs.89626	parathyroid hormone-like hormone			25.20			
_	134806	AD001528	Hs.89718	spermine synthase					4.58	
5	134853	BE268326	Hs.90280	5-aminoimidazole-4-carboxamide ribonucle					4.79	
	134859	D26488	Hs.90315	KIAA0007 protein			6.20			
	134891	R51083	Hs.90787	ESTs			7.40			
	134960	BE246400	Hs.285176	acetyl-Coenzyme A transporter	4.00					
10	134993	BE409809	Hs.301005	purine-rich element binding protein B					4.48	
10	135047	AL134197	Hs.93597	cyclin-dependent kinase 5, regulatory su	9.50					
	135080	AI761180	Hs.94211	rcd1 (required for cell differentiation,	5.00					
	135103	NM_003428	Hs.9450	zinc finger protein 84 (HPF2)		11.00				
	135145	AW014729	Hs.95262	nuclear factor related to kappa 8 bindin			71.00		4.01	
15	135184	U13222	Hs.96028	forkhead box D1	40.50		7.00			
13	135242	AI583187	Hs.9700	cyclin E1	13.50					
	135286	AW023482	Hs.97849	ESTs	6.46	8.80				
	135289	AW372569	Hs.9788	hypothetical protein MGC10924 similar to ATP-dependent RNA helicase	10.00	0.00				
	135355	AK001652	Hs.99423 Hs.997	protease, serine, 22	8.00					
20	135371 135393	NM_006025 L11244	Hs.99886	complement component 4-binding protein,	0.00			14.60		
20	100080	L11244	113.33000	complement component volticing protein,				17,00		

TABLE 5B shows the accession numbers for those primekeys lacking unigenelD's for Table 5A. For each probeset we have listed the gene cluster number from which the oligonucleotides were designed. Gene clusters were compiled using sequences derived from Genbank ESTs and mRNAs. These sequences were clustered based on sequence similarity using Clustering and Alignment Tools (DoubleTwist, Oakland California). The Genbank accession numbers for sequences comprising each cluster are listed in the "Accession" column.

	Pkey:	Unique Eos prob	peset identifier number
	CAT numb	er: Gene cluster nu	mber
30	Accession:	Genbank access	sion numbers
	Pkey	CAT number	Accessions
			•
25	117079	1621717_1	H92325 T97125
35	124305	242183_1	AW963221 AA344870 AA344871 H93331
	101502	182026	M26958
	109792	754958_1	R49625 F10674
	126034	1598157_1	H60340 N91637
40	102768	44641_1	U82321 H66077
40	126345	1653833_1	N49713 N49819 W03810
	127066	1703458_1	R25066 R20144 R20145 Z43845
	127099	244301_1	AA347668 AW956810 Z44271 F07065 F07084 R13506
	119243	1774795_1	T12603 T12604
15	125875	1566433_1	H14480 N98295
45	112054	1538292_1	R43590 F10439
	126979	171411_1	AA210954 AA211007
	126992	880655_1	AI809521 H12174 Z42556
	122318	292419_1	AA429743 AA442754
50	114699	135322_1	AA127386 R15644 AA127404
50	114793	150742_1	AA158245 AA158235
	108305	111550_1	AA071391 AA069892 AA069891
	108393	113411_1	AA075211 AA076245 AA075126 AA074946
	100867	tigr_HT4586	U14622
55	123731	genbank_AA6098	39 AA609839
22	109700	genbank_F09609	
	120715	genbank_AA29270	
	113702	genbank_T97307	
	115113	genbank_AA2564	
60	101045	entrez_J05614	J05614
OU	108554	genbank_AA08494	
	108573	genbank_AA0860	
	119052	149538_1	R10889 R10889
	126522	416020_1	W31912 A1167491
65	126605	439280_1	AA676910 AA778853 AA778865 W86800 W42667 AJ580740 AJ690440 AJ561350 AW467906 AW151450 AJ825927 AL041716 AJ885600 AJ742213 AW248624 AJ955498 AA033947
05	103768	46922_1	AA845593 Al623711 N68583 C00064 AA193567 AW083868 AW163216 AA191595 AA522778 Al628008 Al915518 AA843508 Al926195
			AA176265 AW167963 AA992115 W93647 AW103502 AW60360 AW103212 AM59513 AA276165 AA024712 AA069988 AA205591 A1591107
			ANTIQUOS AVEITIGOS ANTIGOS ANT
			H44848 H20477 T91695 W47039 AA070055 AA024795 AA328855 AA379248 AA379330 AA385860 W25920 W03688 AA448359 AA093881
70			AW362477 AA089997 AJ360265 W93479 N99688 AA932257 AW361469 H68590 AA663402 AA069771 AW087986 AI858420 AA600214
70			AISTOT74 AIBST712 AIGB3081 AIBB5584 AW131150 AI567981 AW002714 AW389973 AW075495 AW168303 AA953714 AW516881 AI357375
			AJ556663 AW512676 AJ570580 AJ023690 AA448216 AJ078853 AJ422707 AA779516 AW026972 AW130082 AW162307 AW438646 AA709332
			AW192394 A1167350 Al217879 Al129152 AA719509 Al350480 AA663418 Al003634 AW118546 AA180281 AA42833 Al258625 AA888881
			AID38759 AA846723 AI248770 AA993694 AI280335 AI885107 AW518649 AA641563 AA995835 AA582621 AI276744 AA436478 AI017360
75			AI620763 AI859887 N73926 AI076327 AI741615 AI160617 AW172819 AI492005 AA677429 AA996334 AI693771 AI950039 AI245629 AI288515
, ,			AJB6618B T93293 AA173262 AA599779 AJ680092 AW439316 AJ084555 AJZ72672 AJ583507 AW473219 AA738132 AW473283 AJ367492
			AA995410 Al689624 AA206353 Al033095 Al040382 AA873630 Al221074 Al934840 Al418680 AA844306 R94503 AA773520 AA8443169
	•		AA219425 AA629658 AI811719 AW411275 AI590981 W37907 AI591178 AI684051 AA983238 AA669347 AA976239 AA704570 AI628339
			AI884391 AI241580 AI003539 AW176687 AA009650 N34566 AI333493 AI186070 AA070827 AA411683 AI280884 AA872023 AA207255
80			AA021576 N71953 AI885888 AW076039 T15777 AI537673 AW248048 H09554 W93480 W47001 AW079 114 AA063160 AA757453 R60788
50			AJ859431 H20478 AA218882 AA757465 AA100995 AJ864135 AJ934209 AA070503 H47008 AA219646 W61039 W93907 AW385050 W37967
			W78028 AA189007 AA479136 R93650 AA442312 T30287 AA847528 AA180262 AA009649 C03892 AW149454 AA310963 AA219593
			AA069747 R29207 AA094784 AA293615 AA447848 AI984167 N90393 C05097 N56499 AW292351 AW149681 AW473256 AA629322 AI004409
			AW105577 AJ954937 AJ811070 AA902422 AW514437 AA535460 AA916877 AW517122 AA974657 AA975649 AW517130 AW517129 F31737
85			W07688 AA193645 AA378994 AA489273 F32267 W39303 AA021181 N86810 AA406524 AA062553 AA436801 H08985 H15979 N40310

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AA436789 AA232172 AW360778 W25862 R60282 AA436530 AA378894 AA187461 AI940535 AA604210 AA089514 AA360421 N88223 N84281 AA209340 N56174 N88374 AA191088 AW247891 AA249013 AA093111 AA972536 AW298594 AA375893 T12139 W28186 AW243849 AI288629 AA843996 W15260 AI188286 AW248079 R15836

genbank_W45552 W45552 genbank_R59904 R59904 genbank_AA227934 AA227934 entrez_A28102 AA496369 AA496646 119599 5 112382 105264 100071 123315

10

Table 6A shows 99 genes up-regulated nonsmokers with lung cancer relative to smokers with lung cancer. These genes were selected from 59680 probesets on the Eos/Affyrmetrix Hu03 Genechip array. Gene expression data for each probeset obtained from this analysis was expressed as average intensity (AI), a normalized value reflecting the relative level of mRNA expression.

Unique Eos probeset identifier number Exemplar Accession number, Genbank accession number Unigene number 15 Pkey: ExAccn: UnigenelD:

Nilgene Title:
Unigene gene title

Unigene gene title

average of AI for samples from non-smokers with adenocarcinoma divided by the 90th percentile of AI for samples from smokers with adenocarcinoma average of AI for samples from non-smokers with squamous cell carcinoma divided by the 90th percentile of AI for samples from smokers with squamous cell 20

carcinoma	

	Pkey	ExAcon	UnigenelD	Unigene Title	R1	R2
25	100971	BE379727	Hs.83213	fatty acid binding protein 4, adipocyte		3.64
23	101174	L17330	Hs.280	pre-T/NK cell associated protein	15.00	
	101296	Y12490	Hs.85092	thyroid hormone receptor interactor 11		2.46
	101304	AA001021	Hs.6685	thyrold hormone receptor interactor 8		12.00
	101806	AA586894	Hs.112408	S100 calcium-binding protein A7 (psorias		2.68
30	101972	S82472	***************************************	gb:beta -pol=DNA polymerase beta (exon a		2.11
	102274	U30930	Hs.158540	UDP glycosyltransferase 8 (UDP-galactose	7.50	
	102394	NM_003816	Hs.2442	a disintegrin and metalloproteinase doma	7.50	
	102832	U92015		gb:Human clone 143789 defective mariner	13.50	
	103010	X52509	Hs.161640	tyrosine aminotransferase	9.50	
35	103439	X98266		gb:H.sapiens mRNA for ligase like protel		2.50
	103563	L02911	Hs.150402	activin A receptor, type i	9.00	204
	103857	Al076795	Hs.45033	lacrimal proline rich protein	10.50	3.94
	104239	AB002367	Hs.21355	doublecortin and CaM kinase-like 1	13.50	12.66
40	104590	AW373062	Hs.83623	nuclear receptor subfamily 1, group 1, m	16.50	12.00
40	104907	AA055829	Hs.196701	ESTs, Weakly similar to ALU1_HUMAN ALU	10.30	2.17
	106131	BE514788	Hs.296244	SNARE protein ESTs	7.00	
	106672	H47233	Hs.30643 Hs.18282	KIAA1134 protein	11.50	
	106872	T56887 AA156238	Hs.32501	ESTs	11.00	2.38
45	106960 106971	Z43846	Hs.194478	Homo saplens mRNA; cDNA DKFZp434O1572 (f	9.50	
73	107982	AA035375	Hs.57887	ESTs, Weakly similar to KIAA0758 protsi	0.00	2.95
	108562	AA100796	(13.57 057	gb:zm26c06.s1 Stratagene pancreas (93720	16.50	
	108599	AB018549	Hs.69328	MD-2 protein	13.00	
	108663	BE219231	Hs.292653	ESTs, Weakly similar to T26845 hypotheti		2.40
50	109247	AA314907	Hs.85950	ESTs	7.00	
	109630	R44607	Hs.22672	ESTs		5.00
	110193	AI004874	Hs.310764	Homo saplens mRNA; cDNA DKFZp434M082 (fr	12,50	
	110234	H24458	Hs.32085	EST	16.50	
	110644	R94207	Hs.268989	ESTs, Highly similar to type II CALM/AF1	8.00	
55	110886	AW274992	Hs.72249	three-PDZ containing protein similar to	17.00	
	111057	T79639	Hs.14629	ESTs	16.50	
	111950	AF071594	Hs.110457	Wolf-Hirschhorn syndrome candidate 1	11.00	3.00
	112291	R53972	Hs.26026	ESTs ankyrin 3, node of Ranvier (ankyrin G)		2.79
60	112956	Z43784 T23699	Hs.75893 Hs.7246	ESTs		4.50
UU	113009 113060	BE564162	Hs.250820	hypothetical protein FLJ14827	9.79	
	113073	N39342	Hs.103042	microtubule-associated protein 1B	32.50	
	113074	AK001335	Hs.31137	protein tyrosine phosphatase, receptor t		3.82
	113121	T48011	Hs.8764	EST		2.21
65	113125	AA968672	Hs.8929	hypothetical protein FLJ11362	19.50	
	113757	AA703095	Hs.18631	ESTs		2.65
	113848	W52854	Hs.27099	hypothetical protein FLJ23293 similar to	6.00	
	113884	Al333076	Hs.28529	chromosome 12 open reading frame 2		6.00
7 0	113936	W17056	Hs.83623	nuclear receptor subfamily 1, group I, m		4.63
70	114875	AA235609	Hs.236443	Homo sapiens mRNA; cDNA DKFZp564N1063 (7.00
	114987	AA251016	Hs.87808	EST		6.00
	115460	AW958439	Hs.38613	ESTs		2.27
	115722	W91892	Hs.59609	ESTs	9.50	9.00
75	116261	AA481788	Hs.190150	ESTs ESTs, Weakly similar to ALU2_HUMAN ALU	8.50	
13	116830	H61037 AB023179	Hs.70404 Hs.9059	KIAA0962 protein	7.50	
	116970 117178	H98675	Hs.269034	ESTs	7.00	2.68
	117757	AF088019	Hs.46732	EST	7.50	2.00
	118283	AA287747	Hs.173012	ESTs, Weakly similar to A46010 X-linked	16.50	
80	118384	AF217525	Hs.49002	Down syndrome cell adhesion molecule		2.50
50	118657	AJ822106	Hs.49902	ESTs		2.39
	120328	AA923278	Hs.290905	ESTs, Weakly similar to protease [H.sapi		3.50
	120404	AB023230	Hs.96427	KIAA1013 protein	7.00	
	120524	AA261852	Hs.192905	ESTs	6.00	
85	120688	AW207555	Hs.97093	Homo sapiens cDNA: FLJ23004 fis, clone L	17.92	

	we	O 02/086 4	143				PCT/US02/12476
	121558	AA412497		gb:zt95g12.s1 Soares_testis_NHT Homo sap		2.95	
	121676	H56037	Hs.108146	ESTs	10.00		
	121936	AI024600	Hs.98612	ESTs	15.00		
	121938	AA428659	Hs.98610	ESTs	14.00		
5	122177	AA435789	Hs.98833	EST	8.93		
•	123442	AA299652	Hs.111496	Homo saplens cDNA FLJ11643 fis, clone HE	13.04		
	123551	AA608837		gb:af03h12.s1 Soares_testis_NHT Homo sap	11.50		
	123756	AA609971	Hs.112795	EST	11.00		
	123861	AA620840		gb:af89g01.s1 Soares_testis_NHT Homo sap		2.50	
10	124371	N24924	Hs.188601	ESTs	6.50		
	127477	BE328720	Hs.280651	ESTs		4.33	
	127591	Al190540	Hs.131092	ESTs		3.02	
	128252	AA455924	Hs.192228	ESTs	7.00		
	128426	Al265784	Hs.145197	ESTs		2.08	
15	128925	R67419	Hs.21851	Homo sapiens cDNA FLJ12900 fis, clone NT		2.11	
	128945	A1990506	Hs.8077	Homo sapiens mRNA; cDNA DKFZp547E184 (fr	10.00		
	129105-		Hs.108681	Homo sapiens brain tumor associated prot	15.50	4.05	
	129235	AW977238	Hs.126084	KIAA1055 protein	C 70	4.25	
20	129506	AB020684	Hs.11217	KIAA0877 protein	6.50	10.00	
20	129595	U09550	Hs.1154	oviductal glycoprotein 1, 120kD (mucin 9	20.00	10.00	
	130160	AA305688	Hs.267695	UDP-Gal:betaGlcNAc beta 1,3-galactosyltr	11.50		
	130340	D82326	Hs.239106	solute carrier family 3 (cystine, dibasi	17.50		
	131220	AB023194	Hs.300855	KIAA0977 protein fatty acid binding protein 7, brain	6.10		
25	131430	AI879148 NM_006152	Hs.26770	lymphoid-restricted membrane protein	0.10	6.15	
23	132114 132458	AA935315	Hs.48965	Homo sapiens cDNA: FLJ21693 fis, clone C		5.58	
	132647	NM_006927		sialyltransferase 4B (bela-galactosidase	7.50		
	132655	D49372	Hs.54460	small inducible cytokine subfamily A (Cy	****	2.53	
	132682	AI077500	Hs.54900	serologically defined colon cancer antig		2.50	
30	132747	AA345241	Hs.55950	ESTs, Weakly similar to KIAA1330 protein		2.83	•
-	132812	R50333	Hs.92186	Leman colled-coil protein		3.82	
	133337	AF085983	Hs.293676	ESTs		5.00	
	133876	AL134908	Hs.771	phosphorylase, glycogen; liver (Hers dis		3.00	
	134119	AW157837	Hs.79226	fasciculation and elongation protein zet		2.06	
35	134464	AA302983	Hs.239720	CCR4-NOT transcription complex, subunit		2.27	
	134542	M14156	Hs.85112	insulin-like growth factor 1 (somatomed)	′	11.50	
	135002	AA448542	Hs.251677	G antigen 7B	87.00	0.50	
	135305	AA203555	Hs.98288	Homo sapiens cDNA FLJ14903 fis, clone PL		6.50	
40							
40	TADICE	chave the seed	erion numbers	for those primekeys lacking unigenelD's for Table 6A. For	each probeset we h	ave listed the gene	cluster number from which the
	oligonuolo	otidae waa da	rianed Cone o	histors were complied using sequences derived from Ganb.	ank ESTs and mRN	As. These sequen	ces were clustered based on sequence
	similarity ı	isino Clusterino	and Alignment	t Tools (DoubleTwist, Oakland California). The Genbank ac	cession numbers fo	r sequences comp	rising each cluster are listed in the
	"Accession						
45							
	Pkey:		s probeset iden	lifier number			
		er: Gene clust					1
50	Accession	: Genbank a	ccession numb	ers			
50							
	Pkey	CAT numb	er Accessions		٠		
	108562	36375_1	AA100796 A	AF020589 AA074629 AA075946 AA100849 AA085347 AA1	26309 AA079311 A	A079323 AA08527	4
55	103439	35330_1	X98266 N41		•		
	123551	genbank_/		AA608837			
	123861	genbank_A		AA620840			
	102832	entrez_U9		U92015			
60	101972	entrez_S8		S82472			
60	121558	genbank_A	A412497	AA412497			

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Table 7A shows 98 genes down-regulated in non-smokers with lung cancer relative to smokers with lung cancer. These genes were selected from 59680 probesets on the Eos/Affymetrix Hu03 Genechip array. Gene expression data for each probeset obtained from this analysis was expressed as average intensity (AI), a normalized value reflecting the relative level of mRNA expression.

5

Pkey: Unique Eos probeset identifier number

ExAccn: Exemplar Accession number, Genbank accession number

UnigeneiD: Unigene Title: Unigene gene title

R1: 80th percentile of AI for samples from smokers with adenocarcinoma divided by the average of AI for samples from non-smokers with adenocarcinoma.

82: 90th percentile of AI for samples from smokers with squamous cell carcinoma divided by the average of AI for samples from non-smokers with squamous cell carcinoma. 10

carcinoma.

		Carcinon	10.			
	Pkey	ExAcon	UnigenelD	Unigene Title	R1	R2
15						404.40
15	100187	D17793	Hs.78183	aldo-keto reductase family 1, member C3		164.10
	100380 100576	D82343 X00356	Hs.18551 Hs.37058	neuroblastoma (nerve tissue) protein calcitonin/calcitonin-related polypeptid	102.40	77.40
	100971	BE379727	Hs.83213	fatty acid binding protein 4, adipocyte	463.80	
	101046	K01160	110.00210	(NONE)	672.00	•
20	101066	AW970254	Hs.889	Charot-Leyden crystal protein	66.00	
	101175	U82671	Hs.36980	melanoma antigen, family A, 2		77.20
	101497	W05150	Hs.37034	homeo box A5	62.80	
	101663	NM_003528	Hs.2178	H2B histone family, member Q	78.00	
25	101677	NM_000715	Hs.1012	complement component 4-binding protein,	186.20 80.08	
23	101745 101941	M88700 S77583	Hs.150403	dopa decarboxylase (aromatic L-amino aci gb:HERVK10/HUMMTV reverse transcriptase	99.20	
	102125	NM_006456	Hs.288215	sialyltransferase	33.20	103.10
	102242	U27185	Hs.82547	relinoic acid receptor responder (tazaro	67.00	
•	102340	U37055	Hs.278657	macrophage stimulating 1 (hepatocyte gro	71.60	
30	102369	U39840	Hs.299867	hepatocyte nuclear factor 3, alpha		69.70
	102457	NM_001394	Hs.2359	dual specificity phosphatase 4	153,00	05.70
	102669	U71207	Hs.29279	eyes absent (Drosophila) homolog 2		65.70 58.80
	102796 102829	AL079848 NM_006183	Hs.107019 Hs.80962	symplekin; Huntingtin interacting protei neurotensin		268.80
35	103207	X72790	113.00302	gb:Human endogenous retrovirus mRNA for	70.00	200.00
-	103242	X76342	Hs.389	alcohol dehydrogenase 7 (class IV), mu o		212.10
	103260	X78416	Hs.3155	casein, alpha		130.70
	103351	X89211		gb:H.saplens DNA for endogenous retrovir	64.60	
40	104212	AB002298	Hs.173035	KIAA0300 protein	66.80	
40	104252	AF002246	Hs.210863	cell adhesion molecule with homology to	63.80	
	104258 105024	AF007216 AA126311	Hs.5462 Hs.9879	solute carrier family 4, sodium bicarbon ESTs	94.40 68.20	
	106260	Al097144	Hs.5250	ESTs, Weakly similar to ALU1_HUMAN ALU S	05.20	74.60
	106440	AA449563	Hs.151393	glutamate-cysteine ligase, catalytic sub		71.10
45	106566	BE298210		gb:601118016F1 NIH_MGC_17 Homo saplens c	73.20	
	106605	AW772298	Hs.21103	Homo seplens mRNA; cDNA DKFZp564B076 (fr	83.80	
	106614	AA648459	Hs.335951	hypothetical protein AF301222		62.30
	106654	AW075485	Hs.286049	phosphoserine aminotransferase		202.40
50	106999 108700	H93281	Hs.10710 Hs.193540	hypothetical protein FLJ20417 ESTs, Moderately similar to 2109260A B c		89.60 66.40
50	108810	AA121518 AW295647	Hs.71331	hypothetical protein MGC5350		95.50
	108857	AK001468	Hs.62180	anillin (Drosophila Scraps homolog), act		63.40
	109597	AA989362	Hs.293780	ESTs	85.00	
	109591	T65568	Hs.12860	ESTs		58.70
55	109704	A1743880	Hs.12876	ESTs	70.40	60.60
	110942	R63503	Hs.28419	ESTs	76.40 74.60	
	111722 112891	R23924 T03927	Hs.23596 Hs.293147	EST ESTs, Moderately similar to A46010 X-II	64.80	
	112992	AL157425	Hs.133315	Homo sapiens mRNA; cDNA DKFZp761J1324 (f	04.00	76.70
60	113073	N39342	Hs.103042	microtubule-associated protein 1B		120.20
	114251	H15261	Hs.21948	ESTs	127.20	
	115230	AA278300	Hs.124292	Homo sapiens cDNA: FLJ23123 fis, clone L	174.00	
	115291	BE545072	Hs.122579	hypothetical protein FLJ10461	CC 40	91.00
65	115815 115909	AW905328 AW872527	Hs.180842 Hs.59761	ribosomal protein L13 ESTs, Weakly similar to DAP1_HUMAN DEATH	66.40	226.60
05	115965	AA001732	Hs.173233	hypothetical protein FLJ10970	82.80	220.00
	116107	AL133916	Hs.172572	hypothetical protein FLJ20093		361.60
	116552	D20508	Hs.164649	hypothetical protein DKFZp434H247	69.00	
70	116571	D45652		gb:HUMGS02848 Human adult lung 3' direct	64.20	
70	118466	N66741		gb:yz33g08.s1 Morton Fetal Cochlea Homo	04.00	63.50
	120484	AA253170 AA398209	Hs.96473	EST EST	81.60	81.10
	120983 121034	AA390209 AI 389951	Hs.97587 Hs.271623	nucleoporin 50kD		66.20
	121423	AW973352	Hs.290585	ESTs	64.40	5515
75	122553	AA451884	Hs.190121	ESTs		60.40
	122946	A1718702	Hs.308026	major histocompatibility complex, class	188.60	
	123130	AA487200		gb:ab19f02.s1 Stratagene lung (937210) H		80.20
	124472	N52517	Hs.102670	EST	71.00	404.00
80	124526	N62096	Hs.293185	ESTs, Weakly similar to JC7328 amino aci ESTs, Moderately similar to ALU7_HUMAN A		104.90 72.00
30	125489 125731	H49193 R61771	Hs.124984 Hs.26912	ESTs Moderately similar to ALO7_HOWAN A		69.90
	- 125747	NM_002884	Hs.865	RAP1A, member of RAS oncogene family	69.00	
	126020	H79863	Hs.114243	ESTs		62.40
0.5	126547	U47732	Hs.84072	transmembrane 4 superfamily member 3		62.80
85	126966	R38438	Hs.182575	solute carrier family 15 (H+/peptide tra		60.10

	W	O 02/086	443				PCT/US02/12476
	127472	AA761378	Hs.192013	ESTs	70.20		
	127610	AA960867	Hs.150271	ESTs, Highly similar to unnamed protein	64.00		
		AW293496	Hs.180138	ESTs	85.20		
		AI022103	Hs.124511	ESTs	96.60		
5		AW889132	Hs.11916	ribokinase	••	78.90	
<i>-</i>		AA650274	Hs.41296	fibronectin leucine rich transmembrane p		106.90	
		AW160432	Hs.296460	craniofacial development protein 1	66.80		
		AW935187	Hs.170162	KIAA1357 protein	00.00	58.53	
		AB040930	Hs.126085	KIAA1497 protein	64.20	00.00	
10	130090		Hs.132390	zinc finger protein 36 (KOX 18)	63.80		
10			Hs.155223	stanniocalcin 2	00.00	139.60	
		AW067800	Hs.63984	cadherin 13, H-cadherin (heart)		64.60	
		AW890487			64.40	04.00	
		AB040900	Hs.6189	KIAA1467 protein	76.20		
1.5		BE501914	Hs.24654	Homo sapiens cDNA FLJ11640 fis, clone HE			
15		AB014548	Hs.31921	KIAA0648 protein	97.80	74.00	
		AB018324	Hs.42676	KIAA0781 protein		71.00	
		NM_001448	Hs.58367	glypican 4	***	88.40	
	132977	AA093322	Hs.301404	RNA binding motif protein 3	133.20		
~~		L20852	Hs.10018	solute carrier family 20 (phosphate tran		59.30	,
20		Al110684	Hs.7645	fibrinogen, B beta polypeptide	341.00		
	134264	AF149297	Hs.8087	NAG-5 protein		64.30	•
	134265	M83772	Hs.80876	flavin containing monooxygenase 3		232.53	
	134346	X84002	Hs.82037	TATA box binding protein (TBP)-associate	66.00		
	134395	AA456539	Hs.8262	lysosomal-associated membrane protein 2		75.80	
25	135047	AL134197	Hs.93597	cyclin-dependent kinase 5, regulatory su		108.30	
	135056	N75765	Hs.93765	lipoma HMGIC fusion partner	71.40		
	135309	Al564123	Hs.42500	ADP-ribosylation factor-like 5	70.40		
20						1 1 1 11.1.211	
30	TABLE 7	B shows the ac	cession number	ers for those primekeys lacking unigeneits for Tal	ble 7A. For each p	probeset we have listed the g	ene cluster number from which the
	oligonuch	eotides were de	signed. Gene	clusters were compiled using sequences derived	from Genbank ES	is and mRNAs. These sequ	ences were clustered based on sequence
	similarity	using Clustering	g and Alignma	nt Tools (DoubleTwist, Oakland California). The G	enbank accession	i numbers for sequences cor	nprising each cluster are listed in the
	"Accessio	on" column.				•	
0.5							
35	Pkey:	Unique Ec	s probeset ide	ntifier number			
	CAT num	iber: Gene clus	ter number				
	Accession	n: Genbank e	ccession num	bers			
40	Pkey	CAT number	r Accession	S			
40							
	103207	306354	X72790				
	106566	120358_1	BE298210	Al672315 AW086489 BE298417 AA455921 AA9)2537 BE327124 F	R14963 AA085210 AW27427	73 A1333584 A1369742 A1039658
			A1885095	AI476470 AI287650 AI885299 AI985381 AW5926	24 AW340136 A12	66556 AA456390 Al310815 .	AA484951
	116571	genbank_D4	15652	D45652			
45	118466	genbank_N6	6741	N66741			
	101046	entrez_K011	60 K01160				
	101941	entrez_S778					
	103351	entrez_X892					
	123130	genbank_A/		AA487200			
50	-20100	30					

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Table 8A shows 1720 genes either up or down-regulated in lung tumors or chronically diseased lung relative to a broad collection of over 40 distinct normal body tissues. Chronically diseased lung samples represent chronic non-malignant lung diseases such as fibrosis, emphysema, and bronchitis. These genes were selected from 39494 probesets on the Eos/Affymetrix Hu02 Genechip array. Gene expression data for each probeset obtained from this analysis was expressed as average intensity (AI), a normalized value reflecting the relative level of mRNA expression.

5

Pkey: Unique Eos probeset identifier number

EXACCN: Exemptar Accession number, Genbank accession number

UnigeneiD: Unigene number

Unigene Title: Unigene gene title

R1: 70th percentile of Al for lung tumors divided by 90th percentile of Al for normal lung

R2: 70th percentile of Al for chronically diseased lung divided by 90th percentile of Al for normal lung 10

	Pkey	ExAcon	UnigeneiD	Unigene Title	R1	R2
15	300097	Al916973	Hs.213603	ESTs	5.46	4.69
	300117	AW189787	Hs.147474	ESTs	0.58	0.56
	300197	Al686661	Hs.218286	ESTs	4.26	5.44
	300201	Al308300	•	gb:ta90c06.x1 NCI_CGAP_Bm20 Homo saplen	0.62	0.83
	300225	A1989963	Hs.197505	ESTs	1.68	1.75
20	300247	AW274682	Hs.161394	ESTs	1.08	2.28
	300256	Al469095	Hs.298241	Transmembrane protease, serine 3	0.86	1.00
	300337	Al707881	Hs.202090	ESTs	5.80	9.09
	300362	Z42308		gb:HSC0FB121 normalized infant brain cDN	4.18	12.78
25	300374	A1859947	Hs.314158	ESTs	2.99	4.38
25	300387	AW270150	Hs.254516	ESTs	1.50 3.98	2.53 5.25
	300440	Al421541	Hs.146164	ESTs EST, Weakly similar to Z232_HUMAN ZINC F	3.18	6.80
	300441 300449	R10387 A1362967	Hs.307921 Hs.132221	hypothetical protein FLJ12401	0.43	0.62
	300449	AW135830	Hs.233955	hypothetical protein FLJ20401	0.16	0.83
30	300552	X85711	Hs.21838	hypothetical protein FLJ11191	4.10	9.75
20	300627	W27363	1.0.0	gb:ab37d01.r1 Stratagene HeLa cell s3 93	4.60	12.60
	300630	AW118822	Hs.128757	ESTs	2.91	5.86
	300716	Al216113	Hs.126280	hypothetical protein FLJ23393	1.00	0.92
	300738	Al623332	Hs.130541	KIAA1542 protein	1.82	1.71
35	300777	AA235361	Hs.96840	KIAA1527 protein	4.48	8.22
	300790	Al492471	Hs.188270	ESTs	1.29	1.18
	300832	A1688147	Hs.220615	ESTs, Weakly similar to T03829 transcrip	5.51	8.56
	300836	Z44942	Hs.22958	calcium channel alpha2-delta3 subunit	4.90 1.70	6.34 2.81
40	300838	Al582897 AW449802	Hs.192570	hypothetical protein FLJ22028 Homo sapiens cONA FLJ20428 fis, clone KA	4.56	7.91
-1 0	300878 300897	A1890356	Hs.285901 Hs.127804	ESTs, Weakly similar to T17233 hypotheti	2.23	1.58
	300926	AA504860	110.127007	gb:ab03a10.s1 Stratagene fetal retina 93	2.13	3.50
	300960	AI041019	Hs.152454	ESTs	2.74	4.46
	300961	AW204069	Hs.312716	ESTs, Weakly similar to unnamed protein	1.00	1.00
45	300962	AA593373	Hs.293744	ESTs	1.46	1.51
	300967	AA565209	Hs.269439	ESTs	0.39	1.30
	300987	AW450840	Hs.148590	ESTs, Weakly similar to AF208846 1 BM-00	1.49	1.08
	300988	AI927208	Hs.208952	ESTs	0.16	0.37 1.94
50	301050	AW136973	Hs.288516	ESTs, Weakly similar to S69890 mitogen I	3.23 6.76	14.28
30	301098 301157	AA677570 AA729905	Hs.185918 Hs.231916	ESTs ESTs	3.16	8.85
	301162	A1142118	Hs.129004	ESTs	1.68	7.18
	301170	AA737594	Hs.247606	ESTs	4.40	6.42
	301192	AI808751	Hs.121188	ESTs	6.38	11.59
55	301193	AA758115	Hs.128350	ESTs, Weakly similar to JC5423 2-hydroxy	4.35	7.78
	301267	AW297762	Hs.255690	ESTs	1.56	1.61
	301281	AA843986	Hs.190586	ESTs	2.19 0.76	1.78 0.76
	301341	AI819198	Hs.208229	ESTS	1.00	1.81
60	301382 301407	AA912839 AW450466	Hs.163369 Hs.126830	ESTs ESTs	1.48	1.51
00	301452	AA975688	Hs.159955	ESTs	0.51	1.46
	301483	AW272467	Hs.254655	Untitled	2.40	5.02
	301494	AI678034	Hs.131099	ESTs	2.79	3.41
	301521	AI733621	Hs.133011	zinc finger protein 117 (HPF9)	0.67	0.67
65	301531	AI077462	Hs.134084	ESTs	2.52	3.76
	301580	AI878959	Hs.73737	splicing factor, arginine/serine-rich 1	7.41	11.92
	301676	Z43570	Hs.27453	ESTs, Moderately similar to G01251 Rar p	8.31	10.70
	301690	F05865	Hs.108323	ubiquitin-conjugating enzyme E2E 2 (homo	2.70 4.20	4.22 8.78
70	301718	F07744	Hs.7987 Hs.286132	DKFZP434F162 protein D15F37 (pseudogene)	5.93	7.04
70	301799 301804	AA3B4252 AA5B1004	Hs.62180	anillin (Drosophila Scraps homolog), act	1.70	0.76
	301822	X17033	Hs.271986	integrin, alpha 2 (CD49B, alpha 2 subuni	1.58	1.36
	301846	R20002	Hs.6823	hypothetical protein FLJ10430	1.00	1.00
	301868	T71508	Hs.13861	ESTs. Weakly similar to pH sensitive max	2.88	5.49
75	301882	T78054		gb:yc97g09.r1 Soares infant brain 1NIB H	2.28	3.80
	301905	Al991127	Hs.117202	ESTs	1.00	1.00
	301948	AA344647	Hs.116724	aldo-keto reductase family 1, member B11	5.28	2.28
	301960	AW070252	Hs.27973	KIAA0874 prolein	5.3B	6.48
90	302011	T91418	Hs.125156	transcriptional adaptor 2 (ADA2, yeast,	3.03	3.42
80	302016	N40834	Hs.23495	hypothetical protein FLJ11252	1.00 0.71	1.25 0.99
	302041	NM_001501		gonadotropin-releasing hormone 2 paired box gene 9	1.60	1.71
	302072 302094	AJ238381 Al286176	Hs.132576 Hs.6786	ESTs	0.52	1.20
	302095	AW044300	Hs.137506	Homo saplens BAC clone RP11-120J2 from 7	2.75	4.93
85	302148	AW269618	Hs.23244	ESTs	3.04	3.87

	W	O 02/080	5443			•
	302155	AI088485	Hs.144759	ESTs	0.45	1.15
	302201	AJ006276	Hs.159003	transient receptor potential channel 6 UDP-Gal;betaGicNAc beta 1,4- galactosylt	0.33 0.52	0.84 0.94
	302202 302206	AF097159 Al937193	Hs.159140 Hs.41143	phosphoinositide-specific phospholipase	2.76	3.65
5	302209	AF047445	Hs.159297	killer cell lectin-like receptor subfami	1.00	1.00
_	302235	AL049987	Hs.166361	Homo sapiens mRNA; cDNA DKFZp564F112 (fr	1.68	1.50
	302290	AL117607	Hs.175563	Homo saplens mRNA; cDNA DKFZp564N0763 (f	1.00	2.11
	302328 302346	AA354849	Hs.23240 Hs.194625	Homo sapiens cDNA FLJ13496 fis, clone PL dynein, cytoplasmic, light intermediate	9.38 3.27	13.08 7.24
10	302340	AL039101 AJ010901	Hs.198267	mucin 4, tracheobronchiai	2.54	1.88
10	302384	Y08982	Hs.202676	synaptonemal complex protein 2	1.00	0.91
	302406	U86751	Hs.211956	CD3-epsilon-associated protein; antisens	2.63	2.67
	302409	AF155156	Hs.218028	adaptor-related protein complex 4, epsil	5.82 3.66	9.34 3.18
15	302423 302432	AB028977 AL080068	Hs.225974 Hs.272534	KIAA1054 protein Homo sapiens mRNA; cDNA DKFZp564J062 (fr	2.44	6.77
13	302432	AF092047	Hs.227277	sine oculis homeobox (Drosophila) homolo	0.44	0.84
	302437	AB024730	Hs.227473	UDP-N-acetylglucosamine:a-1,3-D-mannosid	4.18	5.64
	302455	AA356923	Hs.240770	nuclear cap binding protein subunit 2, 2	1.85 2.04	0.92 2.13
20	302472 302476	AA317451 AF182294	Hs.6335 Hs.241578	SWI/SNF related, matrix associated, acti U6 snRNA-associated Sm-like protein LSm8	1.44	1.89
20	302489	T80660	Hs.230424	Homo sapiens cDNA FLJ13540 fis, clone PL	0.51	1.10
	302490	AA885502	Hs.187032	ESTs	2.64	4.87
	302562	AJ005585	Hs.48956	gap junction protein, beta 6 (connexin 3	5.34	2.68
25	302566	AA085996 AB029488	Hs.248572 Hs.272100	hypothetical protein FLJ22965 SMS3 protein	1.00 0.52	1.21 1.24
23	302630 302634	AB032953	Hs.173560	odd Oz/ten-m homolog 2 (Drosophila, mous	1.00	1.00
	302638	AA463798	Hs.102696	MCT-1 protein	1.58	1.02
	302647	X57723	Hs.198273	NADH dehydrogenase (ubiquinone) 1 beta s	2.72	6.85
30	302655	AJ227892	Hs.146274	ESTs Homo sapiens, clone IMAGE:2823731, mRNA,	1.00 2.97	4.32 0.93
30	302656 302668	AW293005 AA580691	Hs.70704 Hs.180789	S164 protein	0.80	0.95
	302679	H65022	110.11001	gb:yu66g11.r1 Weizmann Olfactory Epithel	1.68	5.04
	302680	AW192334	Hs.38218	ESTs	2.70	7.98
25	302697	AJ001408		gb:Homo saplens mRNA for Immunoglobulin	4.25 3.91	8.13 8.68
35	302705 302711	U09060 L08442		gb:Human immunoglobulin heavy chain, V-r ab:Human autonomously replicating sequen	2.20	2.73
	302719	W69724	Hs.288959	hypothetical protein FLJ20920	0.54	1.02
	302742	L12069		gb:Homo saplens (clone WR4.10VH) anti-th	4.28	11.57
40	302755	AW384815	Hs.149208	KIAA1555 protein	1.57 2.94	2.38 4.68
40	302771 302789	H98476 AJ245067	Hs.42522	ESTs gb:Homo sapiens mRNA for immunoglobulin	3.49	6.31
	302795	AJ245313	Hs.272838	hypothetical protein FLJ10494	0.80	2.74
	302802	Y08250		gb:H.saplens mRNA for variable region of	1.13	0.77
45	302803	AA442824	Hs.293961	ESTs, Moderately similar to putative DNA	3.14 3.04	10.68 8.24
43	302812 302847	N31301 X98940	Hs.152664	hypothetical protein FLJ20051 gb:H.sapiens rearranged ig heavy chain (1.80	1.92
	302885	AL137763	Hs.132127	hypothetical protein LOC57822	1.00	1.00
	302943	AI581344	Hs.127812	ESTs, Weakly similar to T17330 hypotheti	0.53	0.67
50	302977	AW263124	Hs.315111	hypothetical protein FLJ12894	2.45 4.88	2.62 8.61
30	303006 303011	AF078950 AF090405	Hs.24139	Homo saplens cDNA: FLJ23137 fis, clone L gb:Homo saplens clone 2A1 scFV anlibody	1.41	1.86
	303013	F07898	Hs.288968	RAB22A, member RAS oncogene family	1.51	1.19
	303061	AF151882	Hs.27693	peptidylprolyl isomerase (cyclophilin)-l	0.72	0.76
55	303077	AF163305	Hs.146286	gb:H.sapiens T-cell receptor mRNA kinesin family member 13A	1.17 4.08	3.90 6.46
55	303090 303091	AA443259 AF192913	Hs.130683	zinc finger protein 180 (HHZ168)	2.50	4.37
	303094	AF195513	Hs.278953	Pur-gamma	5.38	8.38
	303095	AF202051	Hs.134079	NM23-H8	3.26	4.08
60	303131 303195	AW081061 AA082211	Hs.103180 Hs.233936	DC2 protein myosin, light polypeptide, regulatory, n	2.02 1.32	1.83 3.95
00	303196	AA082298	Hs.59710	ESTs	0.77	0.53
	303216	AA581439	Hs.152328	ESTs	0.24	0.63
	303222	AA333538	Hs.204501	hypothetical protein FLJ10534	3.56 2.28	6.22 3.17
65	303234 303251	AA132255 AW340037	Hs.143951 Hs.115897	ESTs protocadherin 12	0.3B	1.02
05	303295	AA205625	Hs.208067	ESTs	2.30	1.00
	303297	T80072	Hs.13423	Homo saplens clone 24468 mRNA sequence	1.86	4.48
	303316	AF033122	Hs.14125	p53 regulated PA26 nuclear protein	0.10 4.54	0.80 9.65
70	303467 303506	AA398801 AA340605	Hs.323397 Hs.105887	ESTs ESTs, Weakly similar to Homolog of rat Z	0.09	0.04
, 0	303552	AA359799	Hs.224662	ESTs, Weakly similar to unnamed protein	1.00	1.72
	303598	AA382814		gb:EST96097 Testis I Homo sapiens cDNA 5	4.96	9.14
	303637	AF056083	Hs.24879	phosphatidic acid phosphatase type 2C	2.06 1.00	2.02 1.24
75	303655 303756	AA504702 A1738488	Hs.258802 Hs.115838	ATPase, (Na+)/K+ transporting, beta 4 po ESTs	1.08	1.43
	303856	AA968589	Hs.180532	glucose phosphate isomerase	1.76	1.31
	303893	N88597	Hs.113503	karyopherin (Importin) beta 3	2.30	2.57
	303907	AW467774	Hs.171880	polymerase (RNA) II (DNA directed) polyp Homo sapiens cDNA FLJ12363 fis, clone MA	3.10 5.06	5.79 11.86
80	303946 303978	AW474196 AW513315	Hs.306637	gb:xo43c12.x1 NCI_CGAP_Ut1 Homo sapiens	5.14	7.31
	303981	AW513804	Hs.278834	ESTs, Weakly similar to ALU1_HUMAN ALU S	2.83	4.06
	303990	AW515465		gb:xu71a11.x1 NCI_CGAP_Kid8 Homo sapiens	1.15	2.35
	303998 303999	AW516449 AW516611		gb:xl68f05.x1 NCI_CGAP_Ut2 Homo sapiens gb:xp70b11.x1 NCI_CGAP_Ov39 Homo sapiens	2.20 4.85	9.35 6.28
85	304006	AW517947		gb:xt66h02.x1 NCI_CGAP_Ut2 Homo saplens	3.21	4.07
-						

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	304008	AW518198	Hs.3297	ribosomal protein S27a	6.50	11.08
	304009	AW518206	Hs.181165	eukaryotic translation elongation factor	1.88	3.27
	304024	T03036		gb:FB21B7 Fetal brain, Stratagene Horno s gb:FB26F2 Fetal brain, Stratagene Horno s	2.15 5.88	3.55 11.80
5	304026 304028	T03160 T03266		gb:FB7C1 Fetal brain, Stratagene Homo sa	5.59	13.46
	304036	T16855	Hs.244621	ribosomal protein S14	6.55	14.43
	304046	T54803		gb:yb42d06.s1 Stratagene fetal spleen (9	6.18	12.19
	304061	T61521		gb:yb73g01.s1 Stratagene ovary (937217) gb:yc04c12.s1 Stratagene lung (937210) H	2.64 0.53	8.23 1.61
10	304063 304097	T62536 R25376	Hs.177592	ribosomal protein, large, P1	6.49	11.67
	304114	R78946		gb:yi87g02.s1 Soares placenta Nb2HP Homo	2.90	4.18
	304122	H28966		gb:ym31a06.s1 Soares Infant brain 1NIB H	1.00	2.76
	304155	H68696		gb:yr78b06.s1 Soares fetal liver spleen gb:yy82d08.s1 Soares_multiple_sclerosis_	0.79 4.28	1.18 11.34
15	304203 304234	N56929 W81608		gb:zd88h06.s1 Soares_fetal_heart_NbHH19W	6.47	11.03
	304267	AA064862	Hs.73742	ribosomal protein, large, P0	1.34	1.16
	304270	AA069711	Hs.297753	vimentin	3.40	5.40
	304287	AA079286	Hs.78466	proteasome (prosome, macropaln) 26S sub gb:zp38g12.s1 Stratagene muscle 937209 H	2.93 3.98	4.42 10.96
20	304348 304415	AA179868 AA290747	Hs.169476	glyceraldehyde-3-phosphate dehydrogenase	3.32	5.99
	304430	AA347682		gb:EST54044 Fetal heart II Homo sapiens	1.00	1.00
	304456	AA411240		gb:zv26g05.s1 Soares_NhHMPu_S1 Homo sapl	1.42	3.33
	304521	AA464716		gb:zx82c11.s1 Soares overy tumor NbHOT H gb:zx02c05.s1 Soares_total_fetus_Nb2HF8_	2.18 5.38	1.15 14.11
25	304526 304542	AA476427 AA482602	Hs.169476	glyceraldehyde-3-phosphate dehydrogenase	4.16	8.23
	304546	AA486074	Hs.297681	serine (or cysteine) proteinase inhibito	0.55	1.20
	304607	AA513322		gb:nh85e08.s1 NCI_CGAP_Br1.1 Homo saplen	1.95	2.10
	304640	AA524440	Hs.111334	ferritin, light polypeptide ribosomal protein S23	2.10 3.33	2.83 12.62
30	304650 304735	AA527489 AA576453	Hs.3463	gb:nm75h11.s1 NCI_CGAP_Co9 Homo saplens	1.33	0.88
50	304760	AA580401		gb:nn13g09.s1 NCI_CGAP_Co12 Homo saplens	3.68	8.14
	304849	AA588157	Hs.13801	KIAA1685 protein	2.77	3,70
	304917	AA602685	Hs.284136	PRO2047 protein	7.16 2.47	11.01 4.24
35	304921 304966	AA603092 AA613893	Hs.297753 Hs.282435	vimentin ESTs	6.78	11.66
55	304987	AA618044	Hs.300697	Immunoglobulin heavy constant gamma 3 (G	0.90	1.23
	305016	AA626876		gb:zu89h06.s1 Soares_testis_NHT Homo sap	6.46	10.17
	305034	AA630128		gb:ab99c04.s1 Stratagene lung (937210) H gb:nr72a12.s1 NCI_CGAP_Pr24 Homo saplens	1.00 5.68	1.00 11.59
40	305072 305111	AA641012 AA644187	Hs.303405	ESTs	1.48	1.37
	305148	AA654070	1101000 100	gb:nt01g08.s1 NCI_CGAP_Lym3 Homo sapiens	1.76	4.61
	305159	AA659166	Hs.275668	EST, Weakly similar to EF1D_HUMAN ELONG	1.00	2.15
	305190 305232	AA665955 AA670052	Hs.169476	gb:ag57d12.s1 Gessler Wilms tumor Homo s glyceraldehyde-3-phosphate dehydrogenase	5.31 0.78	8.14 1.18
45	305232	AA670032 AA670480	NS. 105470	gb:ag37e01.s1 Jia bone marrow stroma Hom	3.11	8.66
••	305245	AA676695	Hs.81328	nuclear factor of kappa light polypeptid	4.38	7.53
	305312	AA700201	11. 400040	gb:zj44f07.s1 Soares_fetal_liver_spleen_	2.13 1.20	2.66 1.40
	305322 305394	AA701597 AA720942	Hs.163019 Hs.300697	EST immunoglobulin heavy constant gamma 3 (G	1.16	0.68
50	305413	AA724659	113.00000	gb:ai10f08.s1 Soares_parathyroid_tumor_N	5.86	9.87
	305447	AA737856		gb:nx10c08.s1 NCI_CGAP_GC3 Homo sapiens	2.21	2.86
	305478	AA745664	Hs.287445	hypothetical protein FLJ11726	3.36 1.00	6.54 2.02
	305483 305528	AA748030 AA769156	Hs.303512	EST gb:nz12e05.s1 NCI_CGAP_GCB1 Homo saplens	6.44	9.10
55	305612	AA782347	Hs.272572	hemoglobin, alpha 2	0.19	0.79
	305614	AA782866		gb:aj09h02.s1 Soares_parathyroid_tumor_N	1.00	1.00
	305616	AA782884 AA806124	Hs.275865	ribosomal protein S18 gb:oe29a12.s1 NCI_CGAP_Pr25 Homo saplens	7.57 4.78	10.20 12.42
	305637 305639	AA806138		gb:oe29c12.s1 NCI_CGAP_Pr25 Homo saplens	0.89	0.70
60	305650	AA807709		gb:nw31e04.s1 NCI_CGAP_GC80 Homo sapiens4		8.71
	305690	AA813477	11. 70740	gb:ai67a05.s1 Soares_testis_NHT Homo sap	4.91	9.40 0.81
	305726 305728	AA828156 AA828209	Hs.73742	ribosomal protein, large, P0 gb:of34a02.s1 NCI_CGAP_Kid6 Horno sapiens	0.19 5.12	9.29
	305759	AA835353		gb:ak72b06.s1 Barstead spleen HPLRB2 Hom	1.66	4.11
65	305792	AA845256		gb:ak84a08.s1 Barslead spleen HPLRB2 Hom	2.34	4.25
	305864	AA864374	Hs.73742	ribosomal protein, large, P0 gb:oh63h08.s1 NCI_CGAP_Kid5 Homo sapiens	0.30 2.10	1.40 5.21
	305901 305910	AA872968 AA875981		gb:no3no6.81 NCI_CGAP_Rd3 Homo saplens gb:nx21h02.81 NCI_CGAP_GC3 Homo saplens	0.32	1.01
	306015	AA897116		gb:am08b07.s1 Soares_NFL_T_GBC_S1 Homo s1	.56	1.12
70	306017	AA897221	Hs.109058	ribosomal protein S6 kinase, 90kD, polyp	5.21	7.90
	306020	AA897630	Hs.130027	EST gb:ok03g03.s1 Soares_NFL_T_GBC_S1 Homo s	1.96 7.38	6.59 20.69
	306063 306065	AA906316 AA906725		gb:ok78g02.s1 NCI_CGAP_GC4 Homo saplens	7.19	13.48
	306104	AA910956		gb:ok85h11.s1 NCI_CGAP_Kld3 Homo sapiens	6.50	9.13
75	306109	AA911861		gb:og21a07.s1 NCI_CGAP_PNS1 Homo saplens	4.21	5.25
	306148 306242	AA917409 AA932805	Hs.288036	tRNA isopentenylpyrophosphate transferas gb:oo60g04.s1 NCI_CGAP_Lu5 Homo saplens	2.20 2.84	2.70 5.35
	306288	AA932605 AA936900		gb:oi53h05.s1 NCI_CGAP_HN3 Homo sapiens	1.60	1.12
00	306325	AA953072	Hs.210546	Interleukin 21 receptor	1.65	2.26
80	306353	AA961382	Hs.275865	ribosomal protein S18	3.78 4.30	6.32 5.74
	306375 306396	AA968650 AA970223	Hs.276018	EST, Moderately similar to JC4662 ribos gb:op09d05.s1 NCI_CGAP_Kid6 Homo sapiens	4.30 0.95	5.74 2.45
	306428	AA975110	Hs.191228	hypothetical protein FLJ20284	3.19	4.10
0.5	306442	AA976899		gb:gg35e09.s1 NCt CGAP_GC4 Homo sapiens	4.67	7.44
85	306446	AA977348		gb:oq72e12.s1 NCI_CGAP_Kid6 Homo saplens	3.92	6.27

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	306458	AA978186		gb:op33c06.s1 Soares_NFL_T_GBC_S1 Homo s	3.35	5.77
	306467	AA983508	Hs.163593	ribosomal protein L18a	3.72	5.37
	306510	AA988546	13- 070000	gb:gr84d07.s1 NCI_CGAP_Lu5 Homo sapiens	1.00 6.61	1.00 10.91
5	306555 306557	AA994304 AA994530	Hs.276083	EST, Weakly similar to RL23_HUMAN 60S R gb:ou57e08.s1 NCI_CGAP_Br2 Homo saplens	16.20	31.83
_	306572	AA995686		gb:os25c12.s1 NCI_CGAP_Kid5 Homo sapiens	2.51	6.52
	306582	AA996248		gb:os18c10.s1 NCI_CGAP_Kid5 Homo sepiens	1.42	3.13
	306598	AI000320	Hs.169476	glyceraldehyde-3-phosphate dehydrogenase	4.91	8.68
10	306605	A1000497	Hs.119500	ribosomal protein, large P2	1.96 0.11	8.60 0.45
10	306656 306676	AI004024 AI005603	Hs.284136	gb:ou11b07.x1 Soares_NFL_T_GBC_S1 Homo s PRO2047 protein	9.56	17.28
	306686	AI015615	10.201100	gb:ov29f10.x1 Soares_testis_NHT Homo sap	1.86	3.60
	306702	AI022565	Hs.307670	EST	1.47	1.19
15	306728	AI027359	Hs.272572	hemoglobin, alpha 2	1.28 3.91	2.83 5.21
15	306751 306767	A1032589 A1038963	Hs.249118	gb:ow70h12.s1 Soares_fetal_liver_spleen_ ESTs	3.33	6.06
	306892	Al092465	113.243110	gb:ga75h12.x1 Soares_fetal_heart_NbHH19W	3.77	7.46
	306897	A1093967		gb:qa33c06.s1 Soares_NhHMPu_S1 Homo sapi	2.12	2.85
20	306956	AJ125111		gb:am66f03.s1 Barstead spleen HPLRB2 Hom	6.10	10.52 1.56
20	306958 307035	Al125152 Al142774	Hs.119122	gb:am55e09.x1 Johnston frontal cortex Ho ribosomal protein L13a	1.72 2.00	4.70
	307041	Al144243	110.113122	gb:qb85b12.x1 Soares_fetal_heart_NbHH19W	9.12	12.56
	307091	AI167439		gb:ox70h06.s1 Soares_NhHMPu_S1 Homo sapl	4.88	8.52
35	307181	Al189251		gb:qc99g06.x1 Soares_pregnant_uterus_NbH	3.55	6.44
25	307297	AI205798	Hs.111334	ferritin, light polypeptide	2.46 5.64	4.65 10.13
	307317 307327	Al208303 Al214142	Hs.147333 Hs.246381	EST CD68 antigen	3.18	5.15
	307382	AJ223158	Hs.147885	ESTs	2.02	3.73
20	307410	Al241715	Hs.77039	ribosomal protein S3A	0.72	0.48
30	307415	Al242118	Un 470570	gb:qh92b02.x1 Soares_NFL_T_GBC_S1 Homo s	2.38 2.60	3.51 5.44
	307423 307426	A1243206 A1243364	Hs.179573	collegen, type I, alpha 2 gb:qh30g11.x1 Soares_NFL_T_GBC_S1 Homo s	3.18	7.67
	307517	A1275055		gb:ql72d03.x1 Soares_NhHMPu_S1 Homo sapi	1.00	1.00
26	307551	Al281556		gb:qu52f11.x1 NCI_CGAP_Lym6 Homo sapiens	3.40	11.20
35	307561	A1282207		gb:qp65a12.x1 Soares_fetal_lung_NbHL19W	4.74 3.50	15.51 7.19
	307608 307657	Al290295 Al306428	Hs.298262	gb:qm01f02.x1 Soares_NhHMPu_S1 Homo sapi ribosomal protein S19	1.76	2.44
	307691	Al318285	113.230202	gb:tb17b01.x1 NCI_CGAP_Ov37 Homo saplens	1.59	1.31
40	307701	Al318583	Hs.276672	EST, Weakly similar to RL6_HUMAN 60S RI	1.90	2.13
40	307718	A1333406	Hs.83753	small nuclear ribonucleoprotein polypept	0.45	0.99
	307730 307760	Al336092 Al342387		gb:qt43b07.x1 Soares_fetal_lung_NbHL19W gb:qt27f07.x1 Soares_pregnant_uterus_NbH	1.51 1.00	0.99 1.00
	307764	Al342731		gb:qo26a07.x1 NCI_CGAP_Lu5 Homo sapiens	4.52	12.58
4 ~	307783	Al347274		gb:tc05d02.x1 NCI_CGAP_Co16 Homo saplens	1.42	1.00
45	307796	Al350556		gb:qt18f09.x1 NCI_CGAP_GC4 Homo saplens	6.57	9.61
	307807 307808	Al351799 Al351826		gb:qt09d02.x1 NCI_CGAP_GC4 Homo saplens gb:qt09g03.x1 NCI_CGAP_GC4 Homo saplens	3.38 0.33	7.68 0.86
	307820	Al355761		gb:qt94a11.x1 NCI_CGAP_Co14 Homo saplens	7.94	21.57
50	307830	Al358722	Hs.276737	EST, Weakly similar to R5HU22 ribosomal	2.05	3.32
50	307852	AI365541		gb:qz08g05.x1 NCI_CGAP_CLL1 Homo sapiens	3.18 3.13	5.21 4.99
	307902 307997	Al380462 Al434512	Hs.181165	gb:tg02h05.x1 NCI_CGAP_CLL1 Homo sapiens eukaryotic translation elongation factor	1.00	3.01
	308002	AI435240	Hs.283442	EST8	5.86	12.64
~ ~	308011	Al439473		gb:ti60a08.x1 NCI_CGAP_Lym12 Homo sapien	3.79	5.83
55	308023	A1452732	Hs.251577	hemoglobin, alpha 1	0.38 4.36	0.88 6.06
	308041 308059	A1458824 A1468938	Hs.169476 Hs.276877	glyceraldehyde-3-phosphate dehydrogenase EST, Weakly similar to RL10_HUMAN 60S R	1.80	1.98
	308085	AI474135	Hs.181165	eukaryotic translation elongation factor	3.38	4.14
	308101	A1475950	Hs.181165	eukaryotic translation elongation factor	1.30	3.87
60	308106	A1476803	11- 20D444	gb:tj77e12.x1 Soares_NSF_F8_9W_OT_PA_P_S2	2.38 2.70	8.72 3.86
	308122 308154	A1480123 A1500600	Hs.309411	EST gb:tn93d08.x1 NCI_CGAP_Ut2 Homo sapiens	0.66	3.86 1.33
	308171	AI523632	Hs.298766	ESTs, Weakly similar to schlafen4 [M.mu	2.48	4.86
~~	308211	AI557029	Hs.278572	anaplastic lymphoma kinase (KI-1)	2.43	2.14
65	308213	AI557041		gb:PT2.1_12_E04.r tumor2 Homo sapiens cD	3.34 4.61	3.79 4.78
	308216 308219	Al557135 Al557246		gb:PT2.1_13_H06.r tumor2 Homo sapiens cD gb:PT2.1_15_D07.r tumor2 Homo sapiens cD	4.87	7.94
	308271	AI567844	Hs.252259	ribosomal protein S3	2.40	6.35
7 0	308319	AI583983	Hs.181165	eukaryotic translation elongation factor	2.45	3.33
70	308362	A1613519	Hs.105749	KIAA0553 protein	1.24	1.41 4.82
	308413 308450	A1636253 A1660860	Hs.196511 Hs.96840	ESTs KIAA1527 protein	3.16 1.79	2.68
	308464	AI672425	Hs.277117	EST, Moderately similar to 138055 myosi	4.87	8.27
25	308588	Al718299		gb:as51g12.x1 Barstead aorta HPLRB6 Homo	3.90	5.64
75	308599	AI719893	Do 404774	gb:as47d07.x1 Barstead aorta HPLRB6 Homo	3.32	5.12 2.36
	308615 308643	A1738593 A1745040	Hs.101774	hypothetical protein FLJ23045 gb:tr19a12.x1 NCI_CGAP_0v23 Homo sapiens	3.11 3.98	2.36 3.69
	308673	A1745040 A1760864		gb:wi09c10.x1 NCI_CGAP_CLL1 Homo sapiens	0.82	0.99
00	308697	A1767143		gb:wi97a07.x1 NCI_CGAP_Kid12 Homo sapien	2.76	5.59
80	308762	A1807405	Hs.259408	ESTs	3.17 1.00	6.30 1.00
	308778 308782	A1811109 A1811767	Hs.2186	gb:tr04c11.x1 NCI_CGAP_0v23 Homo saplens eukaryotic translation elongation factor	2.94	5.15
	308808	Al818289	100	gb:wk52c01.x1 NCI_CGAP_Pr22 Homo saplens	4.41	8.34
0.5	308823	A1824118	Hs.217493	annexin A2	1.85	1.92
85	308875	AI832332		gb:at48g03.x1 Barstead colon HPLRB7 Homo	2.52	3.80

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	308879	AI832763	Hs.75968	thymosin, beta 4, X chromosome	3.38	7.96
	308886	AI833240	113.70000	ab:at76d10.x1 Barstead colon HPLRB7 Homo	3.06	2.65
	308898	A1858845		gb:wl32d10.x1 NCI_CGAP_Ut1 Homo saptens	2.45	3.44
	308934	A1865023	Hs.177	phosphatidylinositol glycan, class H	4.14	6.76
5	308966	AIB70704		gb:wi47h01.x1 NCI_CGAP_Ut1 Homo sapiens	1.00 7.15	1.00 11.10
	308979	AI873111		gb:wi52h05.x1 NCI_CGAP_Bm25 Homo sapien gb:tq39f01.x1 NCI_CGAP_Ut1 Homo sapiens	0.61	0.59
	309045 309051	Al910902 Al911975		gb:wd78d01.x1 NCI_CGAP_Lu24 Homo sapiens	1.78	4.42
	309069	AI917366	Hs.78202	SWI/SNF related, matrix associated, act	3.27	5.88
10	309083	A1922426	Hs.119598	ribosomal protein L3	2.39	3.34
	309105	A1925503	Hs.265884	ESTs	5.54	17.78 2.92
	309122	Al928178	11- 400040	gb:wo95a11.x1 NCI_CGAP_Kid11 Homo sapien	1.00 1.38	5.55
	309128	AI928816	Hs.180842	ribosomal protein L13 gb:wp84b09.x1 NCI_CGAP_Bm25 Homo saplen	2.43	3.11
15	309164 309177	Al937761 Al951118		gb:wx63g05.x1 NCI_CGAP_Br18 Homo sapiens	0.81	0.97
13	309288	Al991525	Hs.299426	ESTs	4.86	7.46
	309299	AW003478	· · · · · · · · · · · · · · · · · · ·	gb:wq66c06.x1 NCI_CGAP_GC6 Homo sapiens	4.36	9.43
	309303	AW004823		gb:ws93a08.x1 NCI_CGAP_Co3 Homo sapiens	2.88	7.54
00	309411	AW085201	Hs.244144	EST	4.30 2.49	7.14 3.11
20	309437	AW090702	Hs.278242	tubulin, alpha, ubiquitous	2.49	4.55
	309459 309476	AW117645 AW129368	Hs.65114	keratin 18 gb:xe14b05.x1 NCI_CGAP_Ut4 Homo sapiens	2.08	6.60
	309499	AW136325	Hs.279771	Homo sapiens clone PP1596 unknown mRNA	2.82	3.55
	309529	AW150807	Hs.181357	laminin receptor 1 (67kD, ribosomal pro	4.78	3.95
25	309532	AW151119		gb:xg33e10.x1 NCI_CGAP_UI1 Homo sapiens	1.18	4.40
	309626	AW192004	Hs.297681	serine (or cysteine) proteinase inhibit	4.46	12.06
	309641	AW194230	Hs.253100	EST, Moderately similar to GHHU Ig gamm	1.47 5.68	1.39 15.20
	309675	AW205681 AW237221	Hs.253506	EST, Moderately similar to ATPN_HUMAN A laminin receptor 1 (67kD, ribosomal prot	1.00	1.00
30	309693 309695	AW237221	Hs.181357 Hs.295605	mannosidase, alpha, class 2A, member 2	5.45	9.61
50	309700	AW241170	Hs.179661	tubulin, beta polypeptide	1.41	1.25
	309747	AW264889		gb:xq36h02.x1 NCI_CGAP_Lu28 Homo saplens	5.00	8.35
	309769	AW272346		gb:xs13c10.x1 NCI_CGAP_Kid11 Homo sapien	5.76	11.90
25	309782	AW275156	Hs.156110	Immunoglobulin kappa constant	0.42 1.00	0.6 9 4.11
35	309783 309799	AW275401 AW276964	Hs.254798	gb:xp58h01.x1 NCI_CGAP_Ov39 Homo saplens	1.68	1.44
	309866	AW299916		gb:xs44c01.x1 NCI_CGAP_Kid11 Homo sapien	3.02	5.04
	309903	AW339071	Hs.300697	immunoglobulin heavy constant gamma 3 (G	1.05	1.18
	309923	AW340684		gb:hd05g08.x1 Soares_NFL_T_GBC_S1 Homo s	2.30	3.67
40	309928	AW341418		gb:hd08c03.x1 Soares_NFL_T_GBC_S1 Homo s	7.41	13.71
	309931	AW341683		gb:hd13d01.x1 Soares_NFL_T_GBC_S1 Homo s	1.20 4.90	12.70 18.29
	309933	AW341936	Hs.257111	gb:hb73f10.x1 NCI_CGAP_Ut2 Homo sapiens hypothetical protein MGC3265	1.99	3.07
	309964 310002	AW449111 AJ439096	Hs.323079	Homo sapiens mRNA; cDNA DKFZp564P116 (fr	0.20	0.47
45	310096	AW136822	Hs.172824	ESTs, Weakly similar to B48013 proline-r	1.51	1.22
	310098	AI685841	Hs.161354	ESTs	0.31	0.76
	310109	Al203094	Hs.148633	ESTs	2.06	5.83
	310112	AW197233	Hs.147253	ESTS	2.92 1.25	3.55 0.84
50	310115	AI611317 AW195642	Hs.223796 Hs.148901	ESTs ESTs	1.00	2.71
50	310121 310148	AJ206614	Hs.197422	ESTs	9.50	15.31
	310193	AI627653	Hs.147562	ESTs	2.85	4.18
	310255	AW450439	Hs.153378	ESTs	4.26	10.63
	310261	AJ240483	Hs.201217	ESTs	3.28 0.26	4.40
55	310264	AI915771	Hs.74170	metallothionein 1E (functional)	5.43	0.86 8.19
	310275 310282	A1242102 A1243332	Hs.213636 Hs.156055	ESTs ESTs	3.15	8.06
	310290	AW013815	Hs.149103	ESTs	2.19	3.12
	310333	AJ253200	Hs.145402	ESTs	1.17	1.91
60	310346	AI261340	Hs.145517	ESTs	4.81	9.95
	310385	AI263392	Hs.156151	ESTs	5.98	7.79 4.63
	310443	AW119018	Hs.164231	ESTS	2.90 0.85	1.01
	310444 310446	AW196632 AI275715	Hs.252956 Hs.145926	ESTs ESTs	2.18	3.85
65	310468	AI984074	Hs.196398	ESTs	3.39	5.19
••	310477	AI948801	Hs.171073	ESTs	1.00	1.00
	310512	AW275603	Hs.200712	ESTs	3.87	8.12
	310514	AI681145	Hs.160724	ESTS	3.30 0.72	7.33 1.44
70	310524	AW082270	Hs.12496	ESTs, Highly similar to AC004836 1 simil ESTs	3.26	3.46
70	310547 310584	A1302654 A1653007	Hs.208024 Hs.156304	EST8	2.39	4.08
	310608	AI962234	Hs.196102	ESTs	5.60	6.49
	310624	Al341594		gb:Human endogenous retrovirus H proteas	4.91	9.09
75	310636	AI814373	Hs.164175	ESTs .	1.85	1.71
75	310648	AI347863	Hs.156672	ESTs Homo sapiens mRNA full length insert cDN	0.17 5.40	0.69 13.22
	310694	AI654370 AI472124	Hs.157752 Hs.157757	ESTs	4.82	6.27
	310695 310714	A)418446	Hs.157882	ESTS	1.76	3.51
	310722	AI989803	Hs.157289	ESTs	1.14	6.85
80	310756	AI916560	Hs.158707	ESTs	8.46	13.01
	310764	Al376769	Hs.167172	ESTs	4.76	7.37
	310848	AI459554	Hs.161286	ESTs	2.84 1.00	1.96 2.32
	310851 310854	AW291714 AJ421677	Hs.221703 Hs.161332	ESTs ESTs	6.37	7.94
85	310858	AI871000	Hs.161330	ESTs	6.07	9.84

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		Al924558	Hs.161399	ESTs	0.87	0.78
	310875	T47764	Hs.132917	ESTs ESTs, Moderately similar to ALU7_HUMAN A	1.00 7.07	3.63 16.68
	310896 310922	AW157731 AW195634	Hs.270982 Hs.170401	ESTS, Moderately Similar to ALO7_HOMAN A	1.00	1.00
5	310955	AI560210	Hs.263912	ESTs	10.08	17.66
_	310957	AW190974	Hs.196918	ESTs	2.18	3.18
	311000	AI521830	Hs.171050	ESTs	3.06	6.64
	311012 311034	AW298070 Al564023	Hs.241097 Hs.311389	ESTs ESTs, Moderately similar to PT0375 natur	1.23 2.44	3.77 2.09
10	311074	AW290922	Hs.199848	ESTs	6.04	14.19
	311134	A1990849	Hs.196971	ESTs	3.54	6.96
	311174	AW450552	Hs.205457	periaxin	0.65	0.95
	311187	A1638374 A1656040	Hs.224189 Hs.196532	ESTs	2.46 1.10	2.78 2.52
15	311220 311230	A1989808	Hs.197663	ESTs	1.41	1.75
	311236	AI653378	Hs.197674	ESTs	2,18	2.11
	311242	AW016812	Hs.200266	ESTs	0.63	5.11
	311258	AJ671221	Hs.199887	ESTs ESTs, Moderately similar to ALU4_HUMAN A	1.00 2.56	1.41 1.94
20	311277 311294	AW072813 AA826425	Hs.270868 Hs.291829	ESTs	1.04	2.69
	311308	F12664	Hs.49000	ESTs	1.98	6.70
	311351	A1682303	Hs.201274	ESTs	4.77	9.38
	311390 311405	AW392997 AW290961	Hs.202280 Hs.201815	ESTs ESTs	2.80 3.80	6.06 11.66
25	311409	A1698839	113.201010	gb:wd31f02.x1 Soares_NFL_T_GBC_S1 Homo s	3.84	6.94
	311420	Al936291	Hs.209867	ESTs	5.30	12.56
	311443	Al791521	Hs.192206	ESTs	4.39	6.09 1.04
	311467 311479	A1934909 A1933672	Hs.175377 Hs.211399	ESTs ESTs	1.00 2.76	5.81
30	311488	R57390	Hs.301064	arfaptin 1	2.50	5.73
	311495	AW300077	Hs.221358	ESTs .	3.63	6.09
	311511 311534	AW444568	Hs.210303 Hs.243549	ESTs ESTs	2.00 0.31	2.87 1.33
	311537	AW130351 AI805121	Hs.211828	ESTs	3.69	5.85
35	311543	AI681360	Hs.201259	ESTs	1.73	1.34
	311551	AW449774	Hs.296380	POM (POM121 rat homolog) and ZP3 fusion	3.31 1.00	6.12 1.00
	311557 311558	A1819230 Z44432	Hs.211238 Hs.63128	interleukin-1 homolog 1 KIAA1292 protein	2.25	3,41
40	311559	AW008271	Hs.265848	similar to rat myomegalin	2.68	5.90
40	311563	AI922143	Hs.211334	ESTs	2.39	3.32 3.85
	311586 311616	A1827834 AW450675	Hs.211227 Hs.212709	ESTs ESTs	2.47 1.00	1.00
	311621	A1924307	Hs.213464	ESTs	4.16	6.74
15	311635	AI928456	Hs.213081	ESTs	2.17	3.76
45	311668 311672	AW193674 R11807	Hs.240044 Hs.20914	ESTs hypothetical protein FLJ23056	2.60 2.79	3.12 5.18
	311683	AW183738	Hs.232644	ESTs	0.19	0.96
	311700	R49601	Hs.171495	retinoic acid receptor, beta	6.28	8.83
50	311714 311735	AW131785 AW294416	Hs.246831 Hs.144687	ESTs, Weakly similar to CIKG_HUMAN VOLTA Homo saplens cDNA FLJ12981 fis, clone NT	5.00 0.96	8.17 0.72
50	311743	T99079	Hs.191194	ESTs	1.00	1.95
	311783	AJ682478	Hs.13528	hypothetical protein FLJ14054	0.16	0.77
	311785	AI056769	Hs.133512 Hs.14014	ESTs ESTs, Weakly similar to KIAA0973 protein	1.34 8.52	3.97 13.32
55	311799 311819	AA780791 AW265275	Hs.254325	ESTs	3.58	3.91
	311823	AI089422	Hs.131297	ESTs	1.40	1.72
	311877	AA349893	Hs.85339	G protein-coupled receptor 39	0.95	0.91
	311886 311896	AA522738 AW206447	Hs.132554	ESTs gb:UI-H-BI1-afg-g-02-0-UI.s1 NCI_CGAP_Su	0.88 1.66	0.87 1.13
60	311910	N28365	Hs.22579	Homo sapiens clone CDABP0036 mRNA sequen	1.66	2.30
	311923	T60843	Hs.189679	ESTs	0.42	2.63
	311933 311959	A1597963	Hs.118726 Hs.124733	ESTs ESTs	1.88 2.02	3.02 2.33
	311960	T67262 AW440133	Hs.189690	ESTs	3.87	6.62
65	311967	Al382726	Hs.182434	ESTs	5.80	8.14
	311975	AA804374	Hs.272203	Homo sapiens cDNA FLJ20843 fis, clone AD	0.98 0.12	3.26 1.39
	312005 312028	T78450 T78886	Hs.13941 Hs.284450	ESTs ESTs	3.78	4.92
	312046	Al580018	Hs.268591	ESTs	4.11	7.32
70	312056	T83748	Hs.268594	ESTs	2.36	3.08
	312064 312088	AA676713 AW303760	Hs.191155 Hs.13685	ESTs ESTs	3.34 1.60	5.28 1.15
	312093	T91809	Hs.121296	ESTs	0.68	0.85
75	312094	Z78390		gb:HSZ78390 Human fetal brain S. Meier-E	3.05	4.48
75	312097 312118	Al352096	Hs.112180	zinc finger protein 148 (pHZ-52) ESTs	4.52 2.40	9.70 2.60
	312118	T85332 Al052609	Hs.178294 Hs.17631	Homo saplens cDNA FLJ20118 fis, clone CO	2.39	3.53
	312147	T89855	Hs.195648	ESTs	0.67	1.03
80	312175	AA953383	Hs.127554	ESTs ESTs	5.85 2.41	10.60 3.32
30	312179 312201	Al052572 Al928365	Hs.269864 Hs.91139	solute carrier family 1 (neuronal/epithe	0.24	0.89
	312207	H90213	Hs.191330	ESTs	2.20	4.55
	312220 312252	N74613 Al128388	Hs.143655	gb:za55a07.s1 Soares fetal liver spleen ESTs	4.28 1.64	11.13 1.57
85	312304	AA491949	Hs.269392	ESTS	0.12	2.47

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\$\frac{3}{2221}\$ Regist \frac{1}{2233}\$ AB25512 \frac{1}{22333}\$ AB25512 \frac{1}{223333}\$ AB25512 \frac{1}{223332}\$ AB2512 \frac{1}{22333232}\$ AB2512 \frac{1}{2233322}\$ AB2512 \frac{1}{2233322}\$ AB2512 \frac{1}{2233322}\$ AB2512 \frac{1}{2233322}\$ AB2512 \frac{1}{2233322}\$ AB2512 AB2512 \frac{1}{2233322							
\$\frac{3}{312339} Au524394 Hs. 165644 \$\frac{1}{31239} Au524394 Hs. 161647 \$\frac{1}{31239} Au524394 Hs. 161647 \$\frac{1}{312315} Au57568 Hs. 16167 \$\frac{1}{312316} Au57668 Hs. 177407 \$\frac{1}{312316} Au57668 Hs. 177407 \$\frac{1}{312316} Au57668 Hs. 177407 \$\frac{1}{31240} Au51133 Hs. 177407 \$\frac{1}{31240} Au51133 Hs. 173407 \$\frac{1}{31240} Au51133 Hs. 173207 \$\frac{1}{31240} Au51133 Hs. 173207 \$\frac{1}{31240} Au51133 Hs. 173207 \$\frac{1}{31240} Au51230 Hs. 173207 \$\frac{1}{31240} Au5922 Hs							
\$\frac{3}{312339} \frac{3}{3424394} \frac{1}{18.16967} \frac{2}{312376} \frac{3}{342596} \frac{3}{3475688} \frac{1}{347568} \frac{1}{347568} \frac{1}{347568} \frac{1}{347568} \frac{1}{347568} \frac{1}{34768} \frac{1}{34768							
31233 ABF5558 N. 1780F STE 0.08 61.73	5						
100 312396 Re52089 18.17277 STF					ESTS	10.08	
312393 ABSS140 St.							
312437				Hs.172717			
1312440	10						
1312451 R59999	10			Hs.133315			
15 312207 A168177 Hs. 149533 ESTs 5.89 8.24							
15							
312546 AISS228 H. 159426 h	1.5						
312564 H21520	15						
20							
20 312599 Al865073 Hs. 125720 ESTs S.75 S.29							
312845 H52121							
312666 A7240582 He_214678 ESTs C.92 C.93 C.93 A745696 He_203679 ESTs C.94 C.95 C.	20						
2312889							
25 312817 177469 15.20425 251's 25							
25 312946 AW152104 Hs. 2803579 3 12937 Al69007 Hs. 28555, Weakly similar to unnamed protein 4.20 6.23 312920 AW28297 Hs. 131316 ESTs, Weakly similar to T203_HUMAN TRANS 1.19 0.71 312925 NS0868 Hs. 2171695 ESTs, Weakly similar to T203_HUMAN TRANS 1.19 0.71 312936 Al69168 Hs. 121525 ESTs 2.00 4.25 312937 Al40506 Hs. 121525 ESTs 2.00 4.25 312937 Al40506 Hs. 121525 ESTs 2.00 4.25 312930 AA497043 Hs. 116695 ESTs Weakly similar to ALU7_HUMAN ALU S 2.30 4.80 312930 AA497043 Hs. 116695 ESTs 3.112 3.60 312930 AA49710 Hs. 157737 ESTS 2.20 2.30 2.13 31300 Al414712 Hs. 157737 ESTS 2.50 313009 AV282055 Hs. 113357 ESTS 2.50 313091 AV282055 Hs. 113357 ESTS 2.50 313091 AV282055 Hs. 113357 ESTS 2.50 313091 AV282055 Hs. 1135584 ESTs 2.50 313091 AV282055 Hs. 113557 ESTS 2.50 313091 AV28205 Hs. 125520 ESTS 2.50 31305 AV2605 Hs. 125520 ESTS 2.50 31305 A							
312873 Al680071 H. 5.173922 ESTs Weakly similar to unnamed protein 4.0 6.23	25						
312902					ESTs, Weakly similar to unnamed protein		
31	•						
312935 Al691981 Hs.126252 ESTs 1.00 1.73							
312978 N24897 Hs.293119 SETs	30						
312980 AA87043 Ha.116885 ESTs 3.12 3.60 3.60 3.60 3.1284 M25871 Ha.177337 ESTs 2.03 2.13 3.60 3.13029 AA731520 Ha.170504 Ha.170504 Ha.14990 AV283005 Ha.19357 ESTs 2.03 2.13 3.13058 AB651300 Ha.12980 Ha.12980 AV33008 Ha.19357 ESTs ESTs 0.96 1.33 3.13058 AV283015 Ha.12980 ESTs C.25 1.50 AV283015 Ha.125802 ESTs 0.25 1.50 AV283015 Ha.125802 ESTs 0.25 1.50 AV283013 AV449171 Ha.168677 ESTs 2.25 1.50 AV283013 AV449171 Ha.168677 ESTs 3.28 5.55 1.50 AV283015 Ha.127802 ESTs 3.28 5.55 1.50 AV283015 Ha.127802 ESTs 3.28 5.55 AV283015 AV2830169 Ha.12764 ESTs ESTs 3.28 5.56 AV28301 AV2830169 Ha.12764 ESTs ESTs AV28301 AV2830169 Ha.12764 ESTs ESTs AV28301 AV2830169 Ha.12764 ESTs ESTs AV28301 AV2830169 Ha.129683 ESTs ESTs AV28301 Ha.129630 AV2830169 Ha.129683 ESTs ESTs AV28301 Ha.129630 AV2830169 Ha.129683 ESTs AV28301 Ha.129630 AV2830169 Ha.129683 ESTs AV28301 Ha.129630 ESTs ESTs AV28301 Ha.129630 AV2830169 Ha.129630 ESTs ESTs AV28301 Ha.129630 H	50						
312880 AA497043 hs.115685 ESTs 2.03 2.13 313028 A72817 hs.177337 ESTs 2.03 2.13 313029 A731502 hs.150504 ESTs 2.03 2.13 313029 A731502 hs.150504 ESTs 5.52 8.42 313029 A731502 hs.150504 ESTs 0.95 1.39 313039 AW229055 hs.119357 ESTs 6.44 10.73 313058 A651330 hs.135684 ESTs 0.25 1.50 313070 A422023 hs.161338 ESTs 0.25 11.50 313070 A422023 hs.161338 ESTs 0.25 11.50 313097 A1676164 hs.204309 ESTs 3.32 5.06 313130 AW449171 hs.168677 ESTs 0.25 11.50 313130 AW449171 hs.168677 ESTs 0.25 11.50 313210 N74077 hs.204038 hs.132750 ESTs 0.49 1.38 313210 N74077 hs.197043 ESTs 0.49 1.38 313236 AW238169 hs.3513 ESTs 0.49 1.38 313236 AW238169 hs.12176 ESTs 0.49 1.36 313236 AW238169 hs.12176 ESTs 0.49 1.36 313236 AW238169 hs.12176 ESTs 0.49 1.36 313237 A1770008 hs.129583 ESTs 0.40 1.30 0.66 313267 A1770008 hs.129583 ESTs 0.23 1.30 0.66 313267 A177008 hs.129583 ESTs 0.23 1.30 0.66 313267 A177008 hs.129583 ESTs 0.23 1.30 0.66 313267 A178048 hs.129583 ESTs 0.23 1.30 0.66 313267 A178048 hs.129583 ESTs 0.23 1.30 0.66 313399 AW378899 hs.202121 ESTs 0.23 1.30 0.30 0.66 313399 AW378899 hs.202121 ESTs 0.23 1.30 0.30 0.66 313409 A478161 hs.127832 ESTs 0.23 1.30 0.30 0.66 313409 A478161 hs.127832 ESTs 0.23 1.30 0.30 0.66 313417 AA741151 hs.137323 ESTs 0.40 0.23 1.30 0.30 0.60 0.30 0.60 0.30 0.60 0.30 0.60 0.30 0.60 0.30 0.60 0.6							
35 313000 Al147412 Hs. 146657 ESTs			AA497043	Hs.115685			
313029 AA731520 hs.179504 ESTs, Weakly similar to unnamed protein 6.48 13.20 313039 AM19220 hs.1.19357 ESTs, Weakly similar to unnamed protein 6.44 10.73 313056 Al561330 hs.135884 ESTs 1.51 2.04 40 313058 D81016 hs.125382 ESTs 0.25 1.50 313070 Al76164 hs.204339 ESTs 3.72 4.56 313070 Al76164 hs.204339 ESTs 3.72 4.56 313130 N2449171 hs.168577 ESTs 3.28 45 313133 Al240838 hs.132750 ESTs 3.28 5.06 313200 AW449171 hs.198074 ESTs 3.328 5.05 313210 N74077 hs.197043 ESTs 5.36 5.52 313210 N74077 hs.197043 ESTs 5.56 313220 AW328169 hs.23216 ESTs 5.36 5.52 313225 N93466 hs.121764 ESTs, Weakly similar to ALU1_HUMAN ALU S 5.16 8.76 313226 AW328169 hs.129583 ESTs 5.36 5.52 313227 Al20291 hs.202121 ESTs, Weakly similar to testicular takil 0.74 2.06 313227 Al30229 hs.12921 ESTs, Weakly similar to testicular takil 0.74 2.06 313230 AV328291 hs.202121 ESTs, Weakly similar to testicular takil 0.74 2.06 313267 Al770008 hs.159650 ESTs 5.36 5.52 313290 Al785327 hs.202121 ESTs, Weakly similar to env prolein [H.s 2.00 4.32 2.31 31325 Al27888 hs.1920141 ESTs Weakly similar to env prolein [H.s 2.00 4.32 2.31 31339 AW37889 hs.194078 ESTs 5.36 5.56 5.56 313414 Al241540 hs.127832 ESTs 5.36 5.56 5.56 313414 Al241540 hs.127832 ESTs 5.36 5.56 5.56 313414 Al241540 hs.132933 ESTs 5.56 5.56 5.56 313414 Al241540 hs.132933 ESTs 5.56 5.56 5.56 5.56 5.56 5.56 5.56 5.5	25						
1909	33						
130.56							
40 313056 Biblot 5 125382 ESTs							
313077 Al422023 Ha.161338 ESTs S.72 4.56							
313097 Al676164 Hs.204339 EST6 3.72 4.56	40						
1313130 AW449171 Hs.168677 ESTs 3.28 5.06							
45 313156 NS9284 Hs.289010 ESTs 0.49 1.36 5.52 313153 Al240838 Hs.132750 ESTs 0.30 0.66 313210 N74077 Hs.197043 ESTs 0.30 0.66 313236 AW238169 Hs.83513 ESTs, Weakly similar to ALU1_HUMAN ALU S 5.16 8.76 313265 N93466 Hs.121764 ESTs, Weakly similar to testicular tekti 0.74 2.06 0.33 3275 Al207604 Hs.129583 ESTs 0.23 1.30 313276 Al207604 Hs.159550 ESTs 0.23 1.30 313292 Al362991 Hs.202121 ESTs, Weakly similar to testicular tekti 0.74 2.06 6.68 9.57 313292 Al362991 Hs.202121 ESTs 0.20 1.33 1.30 1.30 1.30 1.30 1.30 1.30 1.3							
A							
Signature Sign	45						
1.00 3.87 313239 W19532 Hs.124170 ESTs, Weakly similar to testicular tekti 0.74 2.06 2.06 2.07 2.06 2.07 2.06 2.07 2.06 2.07 2.06 2.07 2.06 2.07 2.07 2.06 2.07							
ST ST ST ST ST ST ST ST							
Strain S							
313275 Al027604 Hs.159660 ESTs 4.02 4.32 4.362991 Hs.205121 ESTs, Weakly similar to env protein (H.s	50						
313290	50						
State			AI753247				
55 313357 AW074848 Hs.201501 ESTs 4.02 5.33 313393 Al674685 Hs.200141 ESTs 1.36 2.68 5.26 313319 AW376889 Hs.194097 ESTs 2.58 5.26 313417 AJ241540 Hs.132933 ESTs 6.57 15.07 313417 AA576052 Hs.193223 Homo sapiens cDNA FLJ11646 fts, clone HE 2.78 4.70 313497 AJ261390 Hs.146085 KIAA1345 protein 0.91 2.37 313516 AA029058 Hs.135145 ESTs 0.23 0.70 313550 AA6280517 Hs.118502 ESTs 0.23 0.70 313560 AA7340151 Hs.209312 ESTs 0.23 0.70 313662 AA740151 Hs.104627 Homo saplens cDNA FLJ10158 fts, clone HE 1.00 1.72 313671 W49823 Hs.104613 RP42 homolog 1.00 1.00 313672 AW468891 Hs.12948 ESTs 3.46 <td></td> <td></td> <td></td> <td></td> <td></td> <td></td> <td></td>							
313393	55						
313399	55						
STS						2.58	
State				Hs.132933			
313499 Al261390 Hs.146085 KIAA1345 protein 0.91 2.37 313516	60						
313516 AA029058 Hs.135145 ESTs	OU						
133556							
AJS569 AJZ73419 Hs. J36146 hypothetical protein FLJ13984 1.68 1.00						0.23	0.70
13688					hypothetical protein FLJ13984		
313662	65						
Name							
No.							
70 313690 A1493591 Hs.78146 platelet/endothelial cell adhesion molec 0.51 0.97 313711 AA398070 Hs.133471 ESTs 0.18 1.01 313723 AA070412 gbzm68c10.s1 Stratagene neuroepilhelium 1.08 1.03 313726 A1744687 Hs.257806 ESTs 2.13 2.99 313774 AW136836 Hs.144583 ESTs 1.38 1.19 2.13 2.99 313790 AW078569 Hs.177043 ESTs 3.88 5.78 313790 AW078569 Hs.177043 ESTs 0.22 2.06 313832 AW271022 Hs.133294 ESTs 1.15 0.91 313834 AW418779 Hs.114889 ESTs 0.68 3.14 313835 AI538438 Hs.159087 ESTS 5.74 8.88 313852 AW270806 Hs.275002 ESTS 5.74 8.88 313854 AW470806 Hs.275002 ESTs 2.09 4.06 313855 AA731470 Hs.163839 ESTs 3.341 4.09 313871 AW471088 Hs.145950 ESTs 3.41 4.09 313871 AW471088 Hs.145950 ESTS 5.28 6.83 313883 AI949384 ESTS 5.28 6.83 313883 AI949384							
313711	70						0.97
1313726		313711	AA398070	Hs.133471			
75 313774 AW136836 Hs. 144583 ESTs 1.38 1.19 313784 AA910514 Hs. 134905 ESTs 3.88 5.78 313790 AW076569 Hs. 177043 ESTs 0.22 2.06 313832 AW271022 Hs. 133294 ESTs 1.15 0.91 313834 AW418779 Hs. 114889 ESTs 0.68 3.14 313835 AI538438 Hs. 159087 ESTs 5.74 8.88 313852 H18633 Hs. 123641 protein tyrosine phosphatase, receptor t 0.16 1.14 313854 AW470806 Hs. 275002 ESTs 2.09 4.06 313865 AA731470 Hs. 163839 ESTs 3.41 4.09 313871 AW471088 Hs. 145950 ESTS 5.28 6.83 313883 AI949384 ESTS 5.28 6.83							
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80 313852 H18633 Hs. 123641 protein tyrosine phosphatase, receptor t 0.16 1.14 313854 AW470806 Hs. 275002 ESTs 2.09 4.06 313865 AA731470 Hs. 163839 ESTs 3.41 4.09 313871 AW471088 Hs. 145950 ESTs 5.28 6.83 313883 AI949384 phosphatase, receptor t 0.16 1.14 4.09 6.83 6.83 6.83 6.83 6.83 6.83 6.83 6.83							
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313865 AA731470 Hs.163839 ESTs 3.41 4.09 313871 AW471088 Hs.145950 ESTs 5.28 6.83 313883 Al949384 cb:nu76d01.s1 NCl_CGAP_Alv1 Homo saplens 2.90 10.91	JU						
313871 AW471088 Hs.145950 ESTs 5.28 6.83 313883 Al949384 cb:nu76d01.s1 NCl_CGAP_Alv1 Homo saplens 2.90 10.91						3.41	
313883 Al949384 ab:nu76d01.s1 NCI_CGAP_Alv1 Homo saplens 2.90 10.91		313871	AW471088		ESTs	5.28	6.83
OJ 313910 AISBSSSU RS. 103443 NOMO BAPIERS CURA PLUTTOTO IIS, GIUTE RE 1.00 1.00	05			U- 400110	gb:nu76d01.s1 NCI_CGAP_Alv1 Homo saplens		
	03	313915	AIR0A7A0	ns.103443	HOMO SAPISIES COMA PER FIRM 18, GUIR RE	1.00	1.00

PCT/US02/12476

	w	O 02/08	6443			
	313926	AW473830	Hs.171442	ESTs	3.40	4.11
	313948	AW452823	Hs.135268	ESTs	5.77	9.15
	313978	AI870175	Hs.13957	ESTs	0.46	0.75
_	313983	AI829133	Hs.226780	ESTs	4.10	6.40
5	314035 314037	AA164199	Hs.270152	ESTs	5.88 1.00	7.90 3.79
	314037	AW300048 AA166970	Hs.275272 Hs.118748	ESTs ESTs	7.60	11.33
	314067	AW293538	Hs.51743	KIAA1340 protein	1.86	1.21
	314103	AI028477	Hs.132775	ESTs	2.90	5.29
10	314107	AA806113	Hs.189025	ESTs	2.00	1.66
	314113	AA218985	Hs.118854	ESTs	0.91	4.17
	314124	AW118745	Hs.9460	Homo sapiens mRNA; cDNA DKFZp547C244 (fr	2.53	3.32 5.08
	314126	AA226431 AA935633	Hs.194628	gb:nc18b12.s1 NCI_CGAP_Pr1 Homo sapiens ESTs	3.13 2.90	6.35
15	314128 314151	AA236163	Hs.202430	ESTs	4.15	6.45
15	314184	AW081795	Hs.233465	ESTs	3.44	4.65
	314192	AW290975	Hs.118923	ESTs	1.00	1.23
	314244	AL036450	Hs.103238	ESTs	2.88	3.67
20	314253	AA278679	Hs.189510	ESTs	4.98	7.16
20	314262	AW086215	Hs.246096	ESTs	0.38	1.94
	314320 314332	AA811598	Hs.275809 Hs.95612	ESTs ESTs	3.34 2.85	5.66 2.09
	314335	AL037551 AA287443	Hs.142570	Homo sapiens clone 24629 mRNA sequence	4.35	4.78
	314340	AW304350	Hs.130879	ESTs, Moderately similar to putative p15	0.77	0.86
25	314351	AA292275	Hs.193746	ESTs	3.07	3.77
	314376	A1628633	Hs.324679	ESTs	4.10	6.11
	314443	AA827125	Hs.192043	ESTs .	6.20	13.67
	314458	Al217440	Hs.143873	ESTS	0.58	2.49 2.62
30	314466	AA767818	Hs.122707 Hs.125507	ESTs DEAD-box protein	2.53 3.94	5.65
30	314478 314482	AI521173 AL043807	Hs.134182	ESTs	1.30	1.44
	314506	AA833655	Hs.206868	Homo sapiens cDNA FLJ14056 fis, clone HE	3.28	3.47
	314519	R42554	Hs.210862	T-box, brain, 1	3.12	6.16
~~	314529	AL046412	Hs.202151	ESTs	3.43	6.87
35	314546	AW007211	Hs.16131	hypothetical protein FLJ12876	1.38	1.00
	314562	AI564127	Hs.143493	ESTs ESTs	2.29 3.87	5.27 5.75
	314579 314580	AW197442 AW451832	Hs.116998 Hs.255938	ESTs, Moderately similar to KIAA1200 pro	0.10	0.71
	314585	AA918474	Hs.216363	ESTs	1.08	1.40
40	314589	AW384790	Hs.153408	Homo sapiens cDNA FLJ10570 fis, clone NT	1.00	1.00
	314592	AA435761	Hs.192148	ESTs	0.90	2.60
	314603	AA418024	Hs.270670	ESTs	4.56	6.29
	314604	AA946582	Hs.8700	deleted in liver cancer 1	3.42 2.97	3.92 4.55
45	314606 314648	AA418241 AA878419	Hs.188767	ESTs gb:EST391378 MAGE resequences, MAGP Homo		1.36
43	314699	AI038719	Hs.132801	ESTs	3.66	4.97
	314701	Al754634	Hs.131987	ESTs	0.03	0.90
	314710	Al669131	Hs.290989	EST	3.40	7.52
50	314750	A1095005	Hs.135174	ESTs	2.80	6.54
50	314767	AW135412	Hs.164002	ESTs	3.20 1.00	4.26 1.00
	314801 314817	AA481027 Al694139	Hs.109045 Hs.192855	hypothetical protein FLJ10498 ESTs	0.91	0.99
	314835	Al281370	Hs.76064	ribosomal protein L27a	5.75	7.44
	314852	AI903735		gb:MR-BT035-200199-031 BT035 Homo saplen	1.68	4.34
55	314853	AA729232	Hs.153279	ESTs	0.60	1.85
	314940	AW452768	Hs.162045	ESTs	10.10	16.20
	314941	AA515902	Hs.130650	ESTs	0.31	1.02 0.37
	314943 314955	Al476797 AA521382	Hs.184572 Hs.192534	cell division cycle 2, G1 to S and G2 to ESTs	2.18 2.59	3.90
60	314973	AW273128	Hs.300268	ESTs	1.05	1.25
•	315004	AA527941	Hs.325351	EST	5.64	13.63
	315006	Al538613	Hs.298241	Transmembrane protease, serine 3	0.52	1.78
	315033	Al493046	Hs.146133	ESTs	2.46	1.00
65	315035	A1569476	Hs.177135	ESTs	0.34	1.33
65	315056	A1202703	Hs.152414	ESTs ESTs	2.10 1.00	2.64 1.30
	315069 315071	AI821517 AA552690	Hs.105866 Hs.152423	Homo sapiens cDNA: FLJ21274 fls, clone C	1.78	1.00
	315073	AW452948	Hs.257631	ESTs	1.17	1.52
	315078	AA568548	Hs.190616	ESTs	3.00	3.79
70	315080	AA744550	Hs.136345	ESTs	1.00	1.00
	315120	AA564991	Hs.269477	ESTs	0.64	1.44
	315175	AI025842	Hs.152530	ESTs	0.61	1.91
	315193 315196	AI241331 AA972756	Hs.131765 Hs.44898	ESTs Homo sapiens clone TCCCTA00151 mRNA sequ	1.06 0.48	0.97 1.96
75	315196 315200	A1808235	Hs.307686	EST TOO 131 MINUTER SEQUENCES	3.76	9.40
	315254	AI474433	Hs.179556	ESTs	5.37	9.36
	315353	AW452608	Hs.279610	hypothetical protein FLJ10493	1.00	1.30
	315397	AA218940	Hs.137516	fidgetin-like 1	3.38	2.24
90	315403	AW362980	Hs.163924	ESTs	2.04	5.23
80	315431	AA622104	Hs.184838	ESTs gb:qh36f02.x1 Soares_NFL_T_GBC_S1 Homo s	2.36 3.46	8.04 7.64
	315454 315455	Al239473 AW393391	Hs.156919	go:qn:solu2.x1 solales_NFL_1_GBC_S1 Hollio's ESTs	3.78	5.76
	315473	AI681671	Hs.312671	ESTs, Moderately similar to OVCA1	0.89	2.15
o	315483	AW512763	Hs.222024	transcription factor BMAL2	2.32	1.96
85	315526	Al193048	Hs.128685	ESTs	1.67	1.78

	W	O 02/08	6443			
	315530	A1200852	Hs.127780	ESTs	1.05	1.01
	315541	Al168233	Hs.123159	sparm associated antigen 4	0.85	0.56
	315552	AW445034	Hs.256578	ESTs ESTs	1.00 2.66	2.22 2.48
5	315562 315577	AA737415 AW513545	Hs.152826 Hs.17283	hypothetical protein FLJ10890	2.20	2.25
_	315587	Al268399	Hs.140489	ESTs	1.00	1.04
	315589	AW072387	Hs.158258	Homo sapiens mRNA; cDNA DKFZp434B1272 (f	0.14	1.05
	315623	AA364078	Hs.258189	ESTs	7.44	12.56
10	315634	AA837085	Hs.220585	ESTs	0.50 0.43	1.40 1.22
10	315668 315677	AA912347 Al932662	Hs.136585 Hs.164073	ESTs ESTs	0.60	1.39
	315706	AW440742	Hs.155556	hypothetical protein FLJ20202	2.18	3.77
	315707	Al418055	Hs.161160	ESTs	2.88	2.63
1.0	315730	H25899	Hs.201591	ESTs	0.11	0.60
15	315745	AI821759	Hs.191856	ESTs	3.50	7.25
	315791	AA678177	11- 200124	gb:zi15a05.s1 Soares_fetal_liver_spleen_	1.78 4.31	2.63 6.23
	315801 315820	AA827752 A1652022	Hs.266134 Hs.258785	ESTs ESTs	2.35	3.01
	315878	AA683336	Hs.189046	ESTs	2.12	2.64
20	315905	AI821911	Hs.209452	ESTs	1.03	1.97
	315923	Al052789	Hs.133263	ESTs	2.63	5.06
	315954	AW276810	Hs.254859	ESTs, Moderately similar to ALU5_HUMAN A	1.21	0.85
	315978	AA830893	Hs.119769	ESTs Homo sapiens cDNA: FLJ21326 fis, clone C	3.09 2.20	3.41 6.82
25	316001 316011	Al248584 AW516953	Hs.190745 Hs.201372	ESTs	0.35	1.63
23	316012	AA764950	Hs.119898	ESTs	6.56	8.13
	316040	AI983409	Hs.189226	ESTs	5.69	10.69
	316048	Al720759	Hs.224971	ESTs	2.84	10.45
20	316076	AW297895	Hs.116424	ESTs	0.30	1.05 1.43
30	316124 316151	Al308862 Al806016	Hs.167028 Hs.156520	ESTs ESTs	1.00 5.80	9.03
	316187	AW518299	Hs.192253	ESTs	1.20	3.96
	316204	AA731509	Hs.120257	ESTs	4.92	6.94
	316232	AW297853	Hs.251203	ESTs	1.48	1.60
35	316275	Al671041	Hs.292611	ESTs, Moderately similar to ALU1_HUMAN A	5.86	12.14
	316291	AW375974	Hs.156704	ESTs	2.73	2.69
	316303 316344	AA740994	Hs.209609 Hs.120610	ESTs ESTs	1.53 3.66	1.26 8.34
	316346	AA744518 Al028478	Hs.157447	ESTs	3.51	6.69
40	316365	Al627845	Hs.210776	ESTs	2.50	4.33
	316380	Al393378	Hs.164496	ESTs	1.16	2.16
	316470	AA809902	Hs.243813	ESTs	5.40	10.34
	316509	AA767310	Hs.291766	ESTs	2.46	2.89 6.04
45	316514 316519	AA768037 Al929097	Hs.291671	gb:od10c11.s1 NCI_CGAP_GCB1 Homo sapiens	4.70 4.41	9.70
73	316609	AW292520	Hs.122082	ESTs	1.00	2.89
	316633	Al125586	Hs.127955	ESTs	2.61	3.72
	316700	AW172316	Hs.252961	ESTs, Weakly similar to ALU1_HUMAN ALU S	3.46	4.64
50	316711	Al743721	Hs.285316	ESTs, Moderately similar to ALU7_HUMAN A	4.45	6.95 2.40
50	316713	AI090671	Hs.134807 Hs.170673	hypothetical protein FLJ12057 ESTs, Weakly similar to AF126780 1 retin	0.30 0.20	1.45
	316715 316787	AI440266 AW369770	Hs.130351	ESTs	4.05	5.53
	316809	AA825839	Hs.202238	ESTs	2.25	3.82
	316811	AA922060	Hs.132471	ESTs	1.00	1.32
55	316812	AW135045	Hs.232001	ESTs	3.28	4.70
	316818	AA827176	Hs.124316	ESTs	0.67 3.53	1.81 6.00
	316824	AA837416	Hs.124299 Hs.172445	ESTs ESTs	0.72	1.56
	316827 316891	Al380429 AW298119	Hs.202536	ESTs	1.64	2.97
60	316951	AA134365	Hs.57548	ESTs ·	1.45	1.08
	316970	AA860172	Hs.132406	ESTs	1.00	1.53
	316971	AA860212	Hs.170991	ESTs	1.08	1.96
	316990	AA861611	Hs.130643	ESTs hypothetical protein FLJ11350	5.44 3.56	10.04 4.37
65	317001 317008	Al627917 AW051597	Hs.233694 Hs.143707	ESTs	0.69	1.37
05	317051	AA873253	Hs.126233	ESTs	6.18	12.72
	317128	AA971374	Hs.125674	ESTs	1.87	2.66
	317129	H12523	Hs.78521	Homo sapiens cDNA: FLJ21193 fis, clone C	4.12	6.64
70	317137	AW341567	Hs.125710	ESTs	2.82	5.12 2.51
70	317196 317212	A1348258 A1866468	Hs.153412 Hs.148294	ESTs ESTs	1.98 1.86	2.83
	317223	AW297920	Hs.130054	ESTs	0.83	1.57
	317224	D56760	Hs.93029	sparc/osteonectin, cwcv and kazal-like d	2.74	0.86
75	317266	AA906289	Hs.203614	ESTs	1.00	1.00
75	317282	AI807444	Hs.176101	ESTs	2.60	4.21
	317285	AW370882	Hs.222080	ESTs ESTs	1.96 7.16	3.49 8.32
	317302 317304	AA908709 AW449899	Hs.135564 Hs.130184	ESTs ESTs	1.38	2.28
	317320	AA927151	Hs.130452	ESTs	3.58	8.13
80	317413	AW341701	Hs.126622	ESTs	2.08	4.92
	317417	AA918420	Hs.145378	ESTs	3.06	4.79
	317452	AA972965	Hs.135568	ESTs	4.22	9.21 4.15
	317519	A1859895	Hs.126860	ESTS ESTs	1.88 3.12	4.15
85	317521 317529	A1824338 A1916517	Hs.126891 Hs.126865	ESTS	2.73	3.34
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	317570	Al733361	Hs.127122	ESTs	1.00	2.43
	317571	AA938663	Hs.199828	ESTs	5.20	11.95
	317598	AW206035	Hs.192123	ESTs	0.33	1.56
_	317627	Al346110	Hs.132553	ESTs	1.50	1.39
5	317650	A)733310	Hs.127346	ESTs	0.48 4.18	1.46 7.14
	317659 317674	AA961216 AW294909	Hs.127785 Hs.132208	ESTs ESTs	2.92	3.20
	317686	AA969051	Hs.187319	ESTs	1.00	1.01
	317692	Al307659	Hs.174794	ESTs	5.33	9.59
10	317701	AI674774	Hs.128014	ESTs	1.00	1.00
	317711	AI733015	Hs.272189	ESTs	5.13	7.81
	317722	A1733373	Hs.128119	ESTs	2.50	6.03 1.30
	317756	AA973667	Hs.128320	ESTs KIAA0258 gene product	1.59 1.00	2.48
15	317777 317799	Al143525 Al498273	Hs.47313 Hs.128808	EST8	1.78	2.11
13	317803	AA983251	Hs.128899	ESTs	0.80	1.06
	317821	AJ368158	Hs.70983	PTPL1-associated RhoGAP 1	0.17	0.68
	317848	AI820575	Hs.129086	Homo sapiens cDNA FLJ12007 fis, clone HE	5.30	8.16
20	317850	N29974	Hs.152982	hypothetical protein FLJ13117	1.30	2.28
20	317861	AW341064	Hs.129119	ESTs	2.18	5.93
	317865	AI298794	Hs.129130	ESTs deoxyribonuclease II beta	4.48 0.44	8.20 0.99
	317869 317881	AW295184 Al827248	Hs.129142 Hs.224398	Homo sapiens cDNA FLJ11469 fis, clone HE	4.06	2.23
	317890	Al915599	Hs.129225	ESTs	4.68	7.48
25	317899	Al952430	Hs.150614	ESTs, Weakly similar to ALU4_HUMAN ALU S	3.14	3.37
	317986	Al005163	Hs.201378	ESTs, Weakly similar to T12545 hypotheti	0.28	1.66
	318001	AW235697	Hs.130980	ESTs	5.12	9.97
	318016	AI016694	Hs.256921	ESTs	1.86 2.92	4.50 5.22
30	318023 318054	AW243058 AW449270	Hs.131155 Hs.232140	ESTs ESTs	3.92	6.37
50	318068	A1024540	Hs.131574	ESTs	1.21	1.27
	318117	Al208304	Hs.250114	ESTs	0.86	1.17
	318187	Al792585	Hs.133272	ESTs, Weakly similar to ALUC_HUMAN IIII	5.90	6.98
25	318223	Al077540	Hs.134090	ESTs	1.05	0.90
35	318240	Al085377	Hs.143610	ESTs	3.10 0.02	2.40 1.05
	318255 318266	Al082692 Al554341	Hs.134662 Hs.271443	ESTs ESTs	6.12	10.55
	318330	Al093840	Hs.143758	ESTs	4.98	7.90
	318369	Al493501	Hs.170974	ESTs	2.46	5.62
40	318428	Al949409	Hs.194591	ESTs	0.77	0.45
	318458	Al149783	Hs.158438	ESTs	3.54	4.92
	318467	AI151395	Hs.144834	ESTs	4.56 2.08	5.62 4.05
	318473 318476	A1939339 A1693927	Hs.146883 Hs.265165	ESTs ESTs	4.22	8.07
45	318487	Al167877	Hs.143716	ESTs	1.47	1.05
	318488	Al217431	Hs.144709	ESTs	1.40	4.14
	318491	T26477	Hs.22883	ESTs, Weakly similar to ALU8_HUMAN ALU S	1.84	1.90
	318499	T25451		gb:PTHI188 HTCDL1 Homo sapiens cDNA 5'/3	2.58	5.20
50	318537	AA377908	Hs.13254	ESTS	3.26 0.35	4.18 1.07
50	318538 318547	N28625 R20578	Hs.74034 Hs.90431	Homo saplens clone 24651 mRNA sequence ESTs	3.22	4.60
	318552	R18364	Hs.90363	ESTs	4.87	9.06
	318575	R55102	Hs.107761	ESTs, Weakly similar to unnamed protein	1.91	1.98
~ ~	318580	T34571	Hs.49007	poly(A) polymerase alpha	2.74	6.22
55	318587	AA779704	Hs.168830	Homo saplens cDNA FLJ12136 fis, clone MA	0.85	2.46
	318596	AJ470235	Hs.172698	EST apolipoprotein A-II	4.88 4.80	4.93 12.51
	318622 318629	T48325 N25163	Hs.237658 Hs.8861	ESTs	0.39	1.04
	318637	AA243539	Hs.9196	hypothetical protein	1.72	3.57
60	318648	T77141	Hs.184411	albumin	6.27	9.91
	318650	AA393302	Hs.176626	hypothetical protein EDAG-1	3.96	8.84
	318671	AA188823	Hs.299254	Homo sapiens cDNA: FLJ23597 fis, clone L	1.53	0.81 2.19
	318679	T58115	Hs.10336	ESTs Homo sapiens cDNA: FLJ21238 fis, clone C	1.00 3.05	3.18
65	318711 318725	A1936475 A1962487	Hs.101282 Hs.242990	ESTs	1.08	2.46
05	318728	Z30201	Hs.291289	ESTs, Weakly similar to ALU1_HUMAN ALU S	0.77	1.33
	318740	NM_002543		oxidised low density lipoprotein (lectin	0.25	1.49
	318776	R24963	Hs.23766	ESTs	1.00	3.01
70	318784	H0014B	Hs.5181	proliferation-associated 2G4, 38kD	2.70	3.86
70	318816	F07873	Hs.21273	ESTs gb:ym04f10.r1 Soares infant brain 1NIB H	3.90 2.25	7.13 3.56
	318865 318879	H10818 R56332	Hs.18268	adenylate kinase 5	1.78	5.00
	318881	Z43224	Hs.124952	ESTs	4.79	14.13
	318894	F08138	Hs.7387	DKFZP564B116 protein	5.31	7.00
75	318901	AW368520	Hs.301528	L-kynurenine/alpha-aminoadipate aminotra	1.03	0.91
	318925	Z43577	Hs.21470	ESTs	2.23	3.80
	318936	Al219221	Hs.308298	ESTS	1.86 5.84	7.16 9.79
	318982 318986	Z44140 Z44186	Hs.269622	ESTs ESTs, Highly similar to MAON_HUMAN NADP-	1.00	1.00
80	318986 319041	Z44186 Z44720	Hs.169161 Hs.98365	ESTs, Weakly similar to weak similarity	3.38	6.11
-	319103	H05896	Hs.4993	KIAA1313 protein	1.00	1.07
	319170	R13678	Hs.285306	putative selenocysteine lyase	3.79	5.03
	319196	F07953	Hs.16085	putative G-protein coupled receptor	1.00	2.98
95	319199	F07361	Hs.13306	ESTs	3.53 5.87	5.66 7.26
85	319242	F11472	Hs.12839	ESTs	5.87	7.26

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	319263	T65331	Hs.81360	Homo saplens cDNA: FLJ21927 fis, clone H	1.81	1.57
	319267	F11802	Hs.6818	ESTs	1.10	4.72
	319270	R13474	Hs.290263	ESTs	4.80	10.40
_	319279	T65094	Hs.12677	CGI-147 protein	1.50	2.11
5	319282	AA461358	Hs.12876	ESTs	1.00	1.00
	319289	W07304	Hs.79059	transforming growth factor, beta recepto	0.18	0.68
	319291	W86578	Hs.285243	hypothetical protein FLJ22029	0.26	0.62
	319293	F12119	Hs.12583	ESTS	3.13 1.10	4.50 1.00
10	319312	Z45481	11- 005000	gb:HSC2QE041 normalized infant brain cDN	0.16	0.73
10	319370 319391	H54254	Hs.325823 Hs.13911	ESTs, Moderately similar to ALU5_HUMAN A ESTs	1.26	2.43
	319396	R06304 H67130	Hs.301743	ESTs	0.70	0.76
	319398	AA359754	Hs.191196	ESTs	2.45	3.59
	319407	R05329	110.101.00	gb:ye91b04.r1 Soares fetal liver spleen	2.00	3.54
15	319425	T82930		gb:yd39f07.r1 Soares fetal liver spleen	4.28	8.81
	319433	R06050	Hs.191198	ĔSŤs	6.15	14.13
	319437	AA282420	Hs.111991	ESTs, Wealdy similar to Y48A5A.1 [C.eleg	3.26	5.68
	319466	AI809937	Hs.116417	ESTs	1.76	5.65
	319471	R06546	Hs.19717	ESTs	4.29	4.84
20	319480	R06933	Hs.184221	ESTs	1.00	1.00
	319484	T91772		gb:yd52a10.s1 Soares fetal liver spleen	2.81	4.88
	319486	Al382429	Hs.250799	ESTS	2.08 2.80	2.82 4.39
	319508	T99898	Hs.270104	ESTs, Moderately similar to ALU8_HUMAN A	1.55	3.25
25	319523	T69499 R83716	Hs.191184 Hs.14355	ESTs Homo sapiens cDNA FLJ13207 fis, clone NT	1.65	1.19
23	319545 319546	R09692	NS. 14000	gb;yf23b12.r1 Soares fetal liver spleen	5.11	8.54
	319552	AA096106	Hs.20403	ESTs	1.89	3.36
	319582	T82998	Hs.250154	hypothetical protein FLJ12973	3.48	4.82
	319586	D78808	Hs.283683	chromosome 8 open reading frame 4	0.26	0.82
30	319604	R11679	Hs.297753	vimentin	1.68	3.41
	319609	AW247514	Hs.12293	hypothetical protein FLJ21103	3.06	4.24
	319611	H14957		gb:ym19c10.r1 Soares infant brain 1NIB H	2.76	4.24
	319653	AA770183	Hs.173515	uncharacterized hypothalamus protein HT0	2.51	3.55
25	319657	R19897	Hs.106604	ESTs	5.32	7.68
35	319658	R13432	Hs.167481	syntrophin, gamma 1	3.35	5.00 12.55
	319661	H08035	Hs.21398	ESTs, Moderately similar to A Chain A, H	5.18 1.58	1.56
	319682	H06382 R15372	Hs.21400 Hs.22664	ESTs ESTs	1.00	1.22
	319708 319742	T77668	Hs.21162	ESTS	2.48	3.13
40	319748	R18178	Hs.295866	Homo sapiens mRNA; cDNA DKFZp434N1923 (f	3.02	4.85
	319772	R76633	Hs.22646	ESTs	4.36	11.61
	319788	AA321932	Hs.117414	KIAA1320 protein	2.56	3.68
	319805	R92857	Hs.271350	likely ortholog of mouse polydom	4.63	6.56
4.5	319812	N74880	Hs.264330	N-acylsphingosine amidohydrolase (acid c	0.63	1.32
45	319834	AA071267		gb:zm61g01.r1 Stratagene fibroblast (937	0.30	0.94
	319878	T78517	Hs.13941	ESTs	3.99	6.44
	319882	AA258981	Hs.291392	ESTS	5.09 3.24	7.36 3.21
	319912 319935	T77559 H79460	Hs.94109 Hs.271722	Homo sapiens cDNA FLJ13634 fis, clone PL ESTs, Weakly similar to ALU1_HUMAN ALU S	4.40	9.42
50	319944	T79248	Hs.133510	ESTs	3.31	5.39
-	319947	AA160967	Hs.14479	Homo sapiens cDNA FLJ14199 fis, clone NT	2.90	4.95
	319962	H06350	Hs.135056	Human DNA sequence from clone RP5-850E9	1.81	1.57
	320007	AA336314		gb:EST40943 Endometrial tumor Homo saple	3.42	6.29
	320018	T83263		gb:yd40h09.r1 Soares fetal liver spleen	2.77	5.14
55	320030	H63789	Hs.296288	ESTs, Wealdy similar to KIAA0638 protein	4.10	6.69
	320032	A1699772	Hs.292664	ESTs, Weakly similar to A46010 X-linked	3.27	3.27
	320040	AA233671	Hs.87164	hypothetical protein FLJ14001	1.81	1.64
	320047	T86564	Hs.302256	EST Sold for the set for the set of the set	3.38 5.90	7.36 16.73
60	320083	AA074108	Hs.120844	FOXJ2 forkhead factor	2.08	4.47
OO	320096 320099	H58138 AW411307	Hs.117915 Hs.114311	ESTs CDC45 (cell division cycle 45, S.cerevis	1.00	1.00
	320112	T92107	Hs.188489	ESTs	2.27	2.06
	320140	H94179	Hs.119023	SMC2 (structural maintenance of chromoso	1.00	1.00
	320188	AW419200	Hs.172318	ESTs	1.26	1.00
65	320193	AA831259	Hs.17132	ESTs	2.58	6.23
	320195	R62203	Hs.24321	Homo sapiens cDNA FLJ12028 fis, clone HE	2.85	4.53
	320199	R78659	Hs.29792	ESTs	0.40	0.94
	320203	AL049227	Hs.124776	Homo saplens mRNA; cDNA DKFZp564N1116 (f	0.84	1.18
70 .	320219	AA327564	Hs.127011	tubulointerstitlal nephritis antigen	1.00	1.17
70 ·	320220	AF054910	Hs.127111	tektin 2 (testicular)	0.18 5.26	1.09 13.75
	320225	AF058989	Hs.128231 Hs.24683	G antigen, family B, 1 (prostate associa ESTs	1.59	1.93
	320231 320260	H03139 NM_003608		G protein-coupled receptor 65	1.38	4.56
	320267	AL049337	Hs.132571	Homo saplens mRNA; cDNA DKFZp564P016 (fr	1.00	1.92
75	320268	H06019	Hs.151293	Homo sapiens cDNA FLJ10664 fis, clone NT	5.58	5.70
	320322	AF077374	Hs.139322	small proline-rich protein 3	1.41	1.01
	320325	Al167978	Hs.139851	caveolin 2	0.05	0.67
	320330	AF026004	Hs.141660	chloride channel 2	2.17	1.26
90	320339	H10807	Hs.281434	Homo sapiens cDNA FLJ14028 fis, clone HE	1.81	2.32
80	320388	H16065	Hs.31286	ESTS	1.00	3.22
	320402	R22291	Hs.23368	Homo saplens clone FLC0578 PRO2852 mRNA,	1.41 2.31	1.36 3.61
	320413	AA203711	Hs.173269 Hs.124136	ESTs ESTs	11.25	20.78
	320432 320438	R62786 AA253352	Hs.293663	ESTs	2.22	3.49
85	320438	W24548	Hs.5669	ESTs	3.53	8.14
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	320448	Al240233	Hs.80887	v-yes-1 Yamaguchi sarcoma viral related	1.42	3.46
	320451	R26944	Hs.180777	Homo saplens mRNA; cDNA DKFZp564M0264 (f	0.87	0.81
	320484	AA094436	Hs.296267	follistatin-like 1	0.65	1.18
5	320499 320514	R32555 AB007978	Hs.24321 Hs.158278	Homo sapiens cDNA FLJ12028 fis, clone HE KIAA0509 protein	3.44 6.44	7.15 13.62
,	320521	N31464	Hs.24743	hypothetical protein FLJ20171	1.48	1.04
	320526	AW374205	Hs.111314	ESTs	3.66	7.87
	320527	R34672	Hs.324522	ESTs	3.16	5.63
10	320536	AA331732	Hs.137224	ESTs	2.83 1.28	5.83 1.00
10	320556 320564	AF054177 AF056209	Hs.14570 Hs.159396	hypothetical protein FLJ22530 peptidylglycine alpha-amidating monooxyg	1.22	0.81
	320587	Z44524	Hs.167456	Homo sapiens mRNA full length Insert cDN	1.84	2.44
	320635	R54159	Hs.80506	small nuclear ribonucleoprotein polypept	1.00	6.25
4.5	320639	AA243258	Hs.7395	hypothetical protein FLJ23182	2.60	2.30
15	320648	N48521	Hs.26549	Homo sapiens mRNA for KIAA1708 protein,	1.00	1.53
	320651 320664	AA489268 Al904216	Hs.111334 Hs.91251	ferritin, light polypeptide hypothetical protein FLJ11198	0.14 5.02	0.79 8.84
	320676	AA132650	Hs.300511	ESTs	3.63	5.37
	320683	R59291	Hs.26638	ESTs, Weakly similar to unnamed protein	0.37	1.31
20	320689	AA334609	Hs.171929	ESTs, Weakly similar to A54849 collagen	1.27	1.02
	320696	AW135016	Hs.172780	ESTs	3.53	4.60
	320714 320727	Al445591 U96044	Hs.181125	gb:yq04a10.r1 Soares fetal liver spleen irmnunoglobulin lambda locus	1.06 1.35	0.85 1.49
	320771	Al793266	Hs.117176	poly(A)-binding protein, nuclear 1	0.04	0.82
25	320794	AA281993	Hs.91226	ESTs	2.96	4.33
	320822	AF100780	Hs.194679	WNT1 inducible signaling pathway protein	0.10	0.79
	320824	AF120274	Hs.194689	artemin	1.16	1.11
	320830	AJ132445 AA317372	Hs.266416	claudin 14 Homo sapiens mRNA; cDNA DKFZp547C136 (fr	1.06 1.36	1.75 1.47
30	320843 320849	D60031	Hs.34744 Hs.34771	ESTs	5.30	7.49
50	320853	A1473796	Hs.135904	ESTs	1.00	1.00
	320896	AB002155	Hs.271580	uroplakin 1B	5.90	2.55
	320921	R94038	Hs.199538	Inhibin, beta C	2.20	1.17
35	320927	AI205786	Hs.213923	ESTS	0.18 1.67	1.46 2.18
33	320957 320997	AJ878933 H22544	Hs.92023	core histone macroH2A2.2 gb:yn69f11.r1 Soares adult brain N2b5HB5	3.26	3.62
	321045	W88483	Hs.293650	ESTs	2.25	4.55
	321046	H27794	Hs.269055	ESTs	2.69	4.25
40	321052	AW372884	Hs.240770	nuclear cap binding protein subunit 2, 2	2.14	2.56
40	321059	A1092824	Hs.126465 Hs.241411	ESTs Homo sapiens mRNA full length insert cDN	1.69 2.76	0.53 5.20
	321062 321067	R87955 AF131782	Hs.241411	Homo sapiens clone 24941 mRNA sequence	4.79	7.41
	321102	AA018306	115.241400	gb:ze40d08.r1 Soares retina N2b4HR Homo	1.79	4.27
4.5	321130	H43750	Hs.125494	ESTs	1.00	3.14
45	321142	AI817933	Hs.298351	ASPL protein	8.73	15.36
	321155	AA336635 AA700289	Hs.99598	hypothetical protein MGC5338 gb:yu76f11.r1 Soares fetal liver spleen	3.04 4.62	5.03 8.39
	321158 321170	N53742	Hs.172982	ESTs	2.21	4.46
	321199	AW385512		gb:yy56d10.s1 Soares_multiple_sclerosis_	5.69	8.01
50	321206	H54178	Hs.226469	Homo sapiens cDNA FLJ12417 fis, clone MA	4.00	7.32
	321225	AL080073	Hs.251414	Homo sapiens mRNA; cDNA DKFZp564B1462 (f	4.17	4.63
	321236 321244	AW371941 AF068654	Hs.18192	Ser/Arg-related nuclear matrix protein (gb:Homo sapiens isolate AN.1 immunoglobu	1.00 2.18	1.00 9.13
	321270	R83560		gb:yv76c06.s1 Soares fetal liver spleen	3.80	5.26
55	321317	AI937060	Hs.6298	KIAA1151 protein	1.81	1.65
	321318	AB033041	Hs.137507	KIAA1215 protein	1.00	1.00
	321325	AB033100	Hs.300646	KIAA protein (similar to mouse paladin) transcription factor BMAL2	0.44 4. 9 4	0.93 4.93
	321342 321356	AA127984 R93443	Hs.222024 Hs.271770	ESTs	3.10	4.66
60	321418	AI739161	Hs.161075	ESTs	2.28	2.54
	321420	Al368667	Hs.132743	ESTs	1.13	0.97
	321430	U05890	11- 000AF	gb:H.sapiens (DIG3) mRNA for immunoglobu	2.42	3.35
	321453	N50080 X13075	Hs.82845	Homo sapiens cDNA: FLJ21930 fls, clone H gb:Human 2a12 mRNA for kappa-immunoglobu	1.60 0.42	3.11 0.72
65	321467 321468	AA514198	Hs.38540	ESTs	2.46	6.50
••	321491	H70665	Hs.292549	ESTs	1.00	1.25
	321498	AW295517	Hs.255436	EŞTs	3.19	6.24
	321504	W02356	Hs.268980	ESTs	2.28	3.86
70	321510 321513	AA703650 H84972	Hs.255748 Hs.108551	ESTs ESTs	2.14 2.78	3.94 5.37
70	321516	Al382803	Hs.159235	ESTs	3.06	7.19
	321565	AI525773	Hs.266514	hypothetical protein FLJ11342	4.89	7.82
	321577	H84260		gb:ys90g04.r1 Soares retina N2b5HR Homo	1.00	1.73
75	321581	AA019964	Hs.28803	ESTs	4.88 1.00	6.73 2.08
75	321582 321587	AA143755 H95531	Hs.21858	trinucleotide repeat containing 3 gb:ys76e02.r1 Soares retina N2b4HR Homo	2.26	4.52
	321626	AA295430	Hs.96322	hypothetical protein FLJ23560	1.95	3.83
	321628	H87064	Hs.161051	ESTs, Moderately similar to ALU6_HUMAN A	0.47	1.02
οΛ	321642	AW085917	Hs.247084	ESTs	1.52	1.38
80	321669	H95404	Hs.294110	ESTS shipt70o12 of Source, NihHMPu, S1 Homo coni	2.17 4.31	2.45 6.95
	321687 321688	AA625149 H97646	Hs.123158	gb:af70c12.r1 Soares_NhHMPu_S1 Homo sapi Homo sapiens cDNA FLJ12830 fis, clone NT	2.82	3.28
	321693	AA700017	Hs.173737	ras-related C3 botulinum toxin substrate	0.51	1.08
0.5	321700	N55160	Hs.167260	ESTs	4.57	7.46
85	321701	AW390923	Hs.42568	ESTs	1.00	1.00

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	321709	N25847	Hs.108923	RAB38, member RAS oncogene family	1.00	1.00
	321710	N35682	Hs.259743	ESTs	2.97	5.26
	321775	AI694875	Hs.202312	Homo sapiens clone N11 NTera2D1 teratoca	1.00 1.68	1.00 0.45
5	321777 321779	Al637993 N42729	Hs.202312 Hs.163835	Homo sapiens clone N11 NTera2D1 teratoca ESTs	0.90	0.90
,	321829	D81993	Hs.8966	tumor endotheliai marker 8	2.69	3.89
	321846	AA281594	Hs.87902	ESTs	5.11	7.64
	321879	AL109670	Hs.302809	ESTs	6.49	9.58
10	321883	AA426494	Hs.46901	KIAA1462 protein	0.28 0.39	0.95 0.95
10	321899	N55158 AF026944	Hs.29468 Hs.293797	ESTs ESTs	6.20	10.76
	321911 321949	R49202	Hs.181694	EST	4.62	10.51
	321955	A1651866	Hs.195689	ESTs	2.89	5.47
1.5	321956	AL110177	Hs.132882	ESTs	0.32	1.25
15	321987	AL133612	Hs.272759	KIAA1457 protein	1.00 4.00	1.83 6.47
	321991 322002	AL133627 AA328801	Hs.158923 Hs.84522	Homo sapiens mRNA; cDNA DKFZp434K0722 (f ESTs	2.10	3.48
	322035	AL137517	Hs.306201	hypothetical protein DKFZp564O1278	1.00	1.90
	322044	AW340926		gbxy51b10.x1 NCI_CGAP_Lu34.1 Homo sapie	3.20	9.67
20	322057	N92197	Hs.154679	synaptotagmin 1	1.55	1.07
	322060	Al341937	11-040220	gb:qt10e03.x1 NCI_CGAP_GC4 Homo saplens	4.59 2.78	7.68 4.52
	322070 322083	U80769 AF074982	Hs.210322 Hs.226031	Homo sapiens mRNA for KIAA1766 protein, ESTs, Highly similar to KIAA0535 protein	3.10	5.52
	322091	AI819863	Hs.106243	ESTs	1.59	1.75
25	322125	R93901		gb:yq16c12.r1 Soares fetal liver spieen	2.06	5.27
_	322130	R98978	Hs.117767	ESTs	10.12	16.49
	322147	AF085919	Hs.114176	ESTs gb:yr88b03.r1 Soares fetal liver spieen	0.94 4.09	0.64 6.67
	322166 322173	AF085958 H52567		gb:yi85d04.r1 Soares_pineal_gland_N3HPG	3.46	4.85
30	322178	H56535		gb:yt88g03.r1 Soares_pineal_gland_N3HPG	0.44	2.54
-	322179	H92891		gb:yl94c02.s1 Soares_pineal_gland_N3HPG	4.52	7.50
	322186	H67346	Hs.269187	ESTs	0.15	0.98
	322196	W87895	Hs.211516	ESTs ESTs	2.20 3.42	5.04 4.84
35	322212 322221	AF087995 Al890619	Hs.134877 Hs.179662	nucleosome assembly protein 1-like 1	0.82	2.14
55	322277	Al640193	Hs.226389	ESTs	3.62	3.98
	322278	AF086283		gb:zd46f01.r1 Soares_fetal_heart_NbHH19W	1.00	1.00
	322284	Al792140	Hs.49265	ESTs	0.66 0.71	2.76 0.70
40	322288 322320	AL037273 AF086419	Hs.7886	pellino (Drosophila) homolog 1 qb:zd78d03,r1 Soares_fetal_heart_NbHH19W	2.02	2.76
40	322336	AA308526	Hs.76152	decorin	2.92	4.44
	322339	W17348		gb:zb18c07.x5 Soares_fetal_lung_NbHL19W	8.50	11.56
	322366	AW404274	Hs.122492	hypothetical protein	0.61	1.34
45	322372	W25624	Hs.153943	ESTs ESTs, Moderately similar to Osf2 [M.musc	7.37 4.78	12.07 10.50
45	322374 322378	Al394663 AF064819	Hs.122116 Hs.201877	DESC1 protein	1.00	1.00
	322388	AI815730	Hs.247474	hypothetical protein FLJ21032	7.09	8.49
	322416	AA223183	Hs.298442	adaptor-related protein complex 3, mu 1	3.20	5.80
50	322419	AA248987	Hs.14084	ring finger protein 7	1.64 0.83	1.57 1.00
30	322425 322431	W37943 AA069222	Hs.34892 Hs.141892	KIAA1323 protein ESTs	3.96	5.22
	322450	AA040131	Hs.25144	ESTs .	5.18	12.67
	322465	AA137152	Hs.286049	phosphoserine aminotransferase	3.41	2.23
<i>e</i>	322467	AF116826	Hs.180340	putative protein-tyrosine kinase	1.00	1.30 2.03
55	322473	AA744286	Hs.266935 Hs.302213	tRNA selenocysteine associated protein ESTs	1.75 1.00	2.03
	322509 322523	T52172 W80398	Hs.193197	ESTS	2.75	5.49
	322527	AF147359	110.100107	gb:Homo saplens full length insert cDNA	1.25	1.27
	322560	Al916847	Hs.270947	ESTs	4.57	8.81
60	322566	W87285	Hs.269587	ESTs gb:zh69c01,r1 Soares_fetal_liver_spleen_	1.00 4.18	1.42 6.94
	322585 322635	AA837622 AA679084		gb:zh90h08.r1 Soares_fetal_liver_spleen_ gb:zh90h08.r1 Soares_fetal_liver_spleen_	2.40	4.85
	322641	AA007352	Hs.256042	ESTs	2.94	4.64
	322653	AI828854	Hs.258538	striatin, calmodulin-binding protein	0.48	0.38
65	322664	AA011522		gb:zi03g07.r1 Soares_fetal_liver_spleen_	1.92	2.18
	322687	Al110759	U= C0043	gb:AF074666 Human fetal liver cDNA libra potassium voltage-gated channel, shaker-	4.14 3.50	6.75 5.00
	322692 322694	AA018117 Al110872	Hs.60843 Hs.279812	PRO0327 protein	1.80	1.72
	322708	AF113674	Hs.283773	clone FLB1727	1.00	3.43
70	322712	AA021328	Hs.23607	hypothetical protein FLJ11109	3.28	3.86
	322766	AW068805	Hs.288467	Homo sapiens cDNA FLJ12280 fis, clone MA	1.63	1.53
	322770	AA045796 A1608591	Hs.122682 Hs.38991	ESTs S100 calcium-binding protein A2	1.53 12.05	1.06 1.94
	322794 322810	A1962276	Hs.127444	ESTs	4.09	6.90
75	322818	AW043782	Hs.293616	ESTs	1.20	1.63
	322820	Al377755	Hs.120695	ESTs	0.21	1.93
	322872	AA827228	Hs.126943	ESTs Homo sapiens cDNA FLJ14035 fis, clone HE	2.04 5.26	1.63 1.22
	322882 322887	AW248508 Al986306	Hs.279727 Hs.86149	phospholnositol 3-phosphate-binding prot	2.80	2.24
80	322913	A1733737	Hs.68837	ESTs	2.38	6.61
	322926	AIB25940	Hs.211192	ESTs	4.02	5.79
	322929	A1365585	Hs.146246	ESTs	0.30	1.14 1.13
	322968	Al905228	Hs.83484 Hs.212760	SRY (sex determining region Y)-box 4 hypothetical protein FLJ13649	2.06 1.18	2.00
85	322971 322981	C15953 AA493252	Hs.159577	ESTs	2.28	2.61

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	322988	C18727	Hs.171941	ESTs	0.39	2.00
	323003	AI733859	Hs.149089	ESTs	3.28	1.00
	323013	AA134042	Hs.191451	ESTs	3.38	5.68
_	323025	AL157565	Hs.315369	Homo saplens cDNA: FLJ23075 fis, clone L	0.06	1.10
5	323032	AW244073	Hs.145946	ESTs	10.18 1.46	21.27 1.90
	323052 323064	R21124 AL119341	Hs.85573 Hs.49359	Homo saplens DC29 mRNA, complete cds Homo saplens mRNA; cDNA DKFZp547E052 (fr	3.08	5.64
	323098	ALT19341 Al700025	Hs.270471	ESTs	2.31	4.49
	323102	AL119913	Hs.163615	ESTs	5.38	11.64
10	323155	AL135041	1.0.1.000.0	gb:DKFZp762K2310_r1 762 (synonym: hmel2)	2.38	5.56
	323176	AW071648	Hs.82101	pleckstrin homology-like domain, family	1.06	1.41
	323191	AA195600	Hs.301570	ESTs	0.73	1.24
	323225	AA205654	Hs.24790	KIAA1573 protein	5.25	11.95
16	323232	AA148722	Hs.224680	ESTs	0.45	1.35
15	323266	AW003362	Hs.243886	nuclear autoantigenic sperm protein (his	1.71	1.83
	323281	A1697556	Hs.292659	ESTS	1.24 12.68	3.21 15.05
	323283 323314	AA256014	Hs.86682 Hs.191501	Homo sapiens cDNA: FLJ21578 fis, clone C ESTs	4.42	9.61
	323314	AA226310 AL134620	Hs.280175	ESTs	2.98	5.93
20	323334	AL134620	Hs.77273	ras homolog gene family, member A	1.98	3.30
	323338	R74219	Hs.23348	S-phase kinase-associated protein 2 (p45	1.62	1.00
	323348	AA233056	Hs.191518	ESTs	1.00	1.07
	323351	AA704103	Hs.24049	ESTs	1.43	1.68
~~	323359	AA234172	Hs.137418	ESTs	0.34	1.18
25	323360	AA716061	Hs.161719	ESTs	3.01	3.71
	323405	AW139550	Hs.115173	ESTs	1.90	8.81
	323420	A1672386	Hs.263780	ESTs	0.29	1.01 1.92
	323434	AW081455	Hs.120219	ESTS	2.27 0.43	0.80
30	323445	AA253103	Hs.135569	ESTs, Weakly similar to NEUROD [H.sapien	3.19	3.85
30	323449 323492	AA282865 H00978	Hs.284153 Hs.20887	Fanconi anemia, complementation group A hypothetical protein FLI 10392	2.70	3.20
	323501	AA182461	Hs.84520	ESTs	2.04	3.31
	323505	A1652287	115.04020	gb:EST382593 MAGE resequences, MAGK Homo2		3.08
	323515	AA282274	Hs.256083	ESTs	2.69	3.40
35	323541	Al185116	Hs.104613	RP42 homolog	1.20	1.09
	323545	AI814405	Hs.224569	ESTs	1.25	1.55
	323635	R63117	Hs.9691	Homo sapiens cDNA: FLJ23249 fis, clone C	0.27	0.72
	323675	AA984759	Hs.272168	tumor differentially expressed 1	3.70	5.80
40	323678	AL042121	Hs.20880	ESTs	3.33	5.10
40	323691	AA317561	Hs.145599	ESTs	1.00	1.00
	323693	AW297758	Hs.249721	ESTs	2.01	1.54
	323746	AW298611	Hs.12808	MARK	4.11	5.53
	323774	AA329806	Hs.321056	Homo sapiens mRNA; cDNA DKFZp586F1322 (f	2.06 3.42	3.70 8.13
45	323856 323857	AA355264 T18988	Hs.267604 Hs.293668	hypothetical protein FLJ10450 ESTs	5.97	12.51
75	323870	AA341774	Hs.129212	ESTs	3.17	4.52
	323876	AL042492	Hs.147313	ESTs	0.36	1.00
	323885	AA344308	Hs.128427	Homo sapiens BAC clone RP11-335J18 from	2.31	3.33
	323911	AL043212	Hs.92550	ESTs	4.38	5.41
50	323919	AA862973	Hs.220704	ESTs	5.80	10.20
	323972	AI869964	Hs.182906	ESTs	3.10	5.14
	324005	AA610011	Hs.208021	ESTs	5.34	10.07
	324036	A1472078	Hs.303662	ESTs	1.00	5.03
<i>5 5</i>	324055	AA528794	Hs.128644	ESTs	0.86	1.00
55 .	324063	AW292740	Hs.272813	dual oxidase 1	0.45	0.91 5.12
	324072	AA381829	Un 202472	gb:EST94855 Activated T-cells I Homo sap	2.82 2.40	2.52
	324092	AW269931	Hs.202473	Homo sapiens cDNA: FLJ22278 fis, clone H	1.32	4.30
	324095 324129	AW377983 Al381918	Hs.298140 Hs.285833	Homo sapiens cDNA: FLJ22502 fis, clone H Homo sapiens cDNA: FLJ22135 fis, clone H	1.40	1.77
60	324132	AW504860	Hs.288836	hypothetical protein FLJ12673	4.24	6.21
00	324214	AA412395	Hs.225740	ESTs	6.96	10.69
	324227	AA295552	Hs.28631	Homo sapiens cDNA: FLJ22141 fis, clone H	0.81	0.53
	324266	AL047634	Hs.231913	ESTs	2.42	4.05
~~	324275	AA429088	Hs.98523	ESTs	3.62	5.38
65	324281	AL048026	Hs.124675	ESTs, Weakly similar to T14742 hypotheti	0.14	0.70
	324290	AA432032	Hs.304420	ESTs	3.71	4.34
	324303	AL118754		gb:DKFZp761P1910_r1 761 (synonym: hamy2)	0.95	0.91
	324312	Al198841	Hs.128173	ESTs	4.08	5.91
70	324325	AL138153	Hs.300410	ESTs	5.88 0.87	8.25 1.25
70	324338	AL138357	Hs.145078 Hs.99807	regulator of differentiation (In S. pomb ESTs, Weakly similar to unnamed protein	1.28	1.00
	324341 324343	AW197734 AW452016	Hs.293232	ESTS	2.54	3.46
	324371	AA452305	Hs.270319	ESTs	5.85	8.36
	324382	AW502749	Hs.24724	MFH-amplified sequences with leucine-ric	0.76	1.64
75	324384	AA453396	Hs.127656	KIAA1349 protein	2.88	5.69
. •	324385	F28212	Hs.284247	KIAA1491 protein	1.81	1.99
	324388	Al924963	Hs.306206	hypothetical protein FLJ11215	1.00	1.00
	324432	AA464510	Hs.152812	ESTs	2.73	2.17
00	324497	AW152624	Hs.136340	ESTs, Weakly similar to unnamed protein	0.71	1.90
80	324510	Al148353	Hs.287425	Homo sapiens cDNA FLJ11569 fis, clone HE	1.00	1.00
	324580	AA492588	11- 40000	gb:ng99c08.s1 NCI_CGAP_Thy1 Homo sapiens	2.18	3.50
	324582	AA506935	Hs.132036	ESTs, Weakly similar to ALU1_HUMAN ALU S	5.96 2 02	11.36 4.22
	324633	AA572994 AW295832	Hs.325489 Hs.134798	ESTs ESTs, Moderately similar to TTL MOUSE TU	2.92 5.48	11.74
85	324640 324675	AW014734	Hs.157969	ESTs	0.39	0.73
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15							5.51
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225187 A653682 Hs.197812 ESTs	20						
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	327085	2	2.50 12.57
	327130		.38 8.04
	327156 327220		1,74 6.58 1,28 1.54
5	327224		5.56 12.91
_	327288	2	2.61 5.40
	327321		2.42 3.11
	327332 327361		i.62 10.58 2.69 4.41
10	327377		2.04 6.72
	327396	2	2.61 4.50
	327414		.00 8.01 i.91 9.65
	327442 327467		5.58 18.01
15	327473	3	3.79 7.48
	327483		3.08 8.87
	327562 327568		0,68 2.86 .00 2.00
	327606		.06 3.61
20	327611	5	5.90 14.26
	327642		.06 8.74 .05 2.08
	327654 327734		.05 2.08 .00 1.00
	327775		.46 11.79
25	327796		.47 5.65
	327840		3.26 6.64 5.84 15.58
	327940 327984		36 1.50
•	328004	· ·	.87 1.42
30	328021		0.59
	328068 328100		2.83 4.68 3.04 5.39
	328101		.54 5.20
25	328113	C	.72 0.91
35	328157	5	5.58 5.16
	328196 328197	, 'S S	5,76 11.13 5,98 10.58
	328264		.11 . 4.88
40	328299		.20 3.06
40	· 328342 328365		.49 1.94 .00 1.00
	328369		.40 7.36
	328381	1	.86 4.93
45	328451	5	i.51 7.56 i.13 0.72
72	328481 328500		.71 3.97
	328530	· 5	.41 7.62
	328600		1.14 10.68
50	328608 328616		.56 8.17 .24 11.91
•	328623	3	.04 5.46
	328632		.70 1.19
	328664 328666		.48 6.80 0.42 26.47
55	328698		.68 14.56
	328700	2	.74 10.22
	328708		.15 0.57 .23 8.91
	328735 328743		.62 6.54
60	328806	0	.22 0.78
	328861	3	.68 10.54 .42 16.36
	328908 328933	5	.42 16.36 .02 5.29
	328934		.73 4.45
65	328949	3	.34 5.41
	329005 329011		.88 7.26 .52 3.72
	329033		.00 1.03
	329037	5	.07 8.16
70	329067	1	.98 2.41
	329134 329157	2	.24 3.25 .30 11.04
	329178		.64 5.02
75	329192	6	.41 15.27
75	329194		.31 0.79 .60 3.75
	329204 329224		.50 3.75 .99 6.11
	329228	0	.83 0.83
90	329288	0	.63 1.01
80	329337 329541	1	.00 1.00 .76 1.68
	329560		.34 2.02
	329588	1	.68 2.22
85	329643 329703		.18 11.77 .00 1.00
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	W	O 02/086	6443			
	329764				5.78	15.50
	329816			•	2.09 3.13	5.44 10.77
	329860 329993				7.83	14.21
5	330020				5.58	13.12
	330036				3.32 4.31	5.57 7.97
	330052 330085			•	1.34	1.76
	330088				4.70	12.46
10	330093				0.44 3.47	1.06 4.83
	330100 330106				2.14	3.61
	330107				3.17	6.87
1.5	330120				5.61	11.89
15	330123 330208				4.50 1.55	12.74 7.62
	330263				13.10	23.38
	330300				2.81	4.98
20	330313 330366			•	3.00 0.67	4.41 0.76
20	330372				4.76	11.82
	330385	AA449749	Hs.182971	karyopherin alpha 5 (Importin alpha 6)	2.14	2.15
	330397 330468	D14659 L10343	Hs.154387 Hs.112341	KIAA0103 gene product protease inhibitor 3, skin-derived (SKAL	0.40 1.11	1.15 0.94
25	330472	L24203	Hs.82237	ataxia-telangiectasia group D-associated	1.67	1.17
	330478	L38486	Hs.296049	microfibrillar-associated protein 4	0.46	1.07
	330493	M27826	Hs.267319 Hs.71642	endogenous retroviral protease guanine nucleotide binding protein (G pr	1.07 0.97	0.95 0.96
	330495 330506	M31328 M61906	Hs.6241	phosphoinositide-3-kinase, regulatory su	0.17	3.66
30	330512	M80563	Hs.81256	S100 calcium-binding protein A4 (calcium	0.60	1.06
	330537	U19765	Hs.2110	zinc finger protein 9 (a cellular retrov	2.81 3.91	2.07 1.49
	330547 330551	U32989 U39840	Hs.183671 Hs.299867	tryptophan 2,3-dioxygenase hepatocyte nuclear factor 3, alpha	1.15	1.03
0.5	330568	U56244		(NONE)	2.83	4.79
35	330599	U90437	U- 0204E	gb:Human RP1 homolog mRNA, 3'UTR region Homo sapiens cDNA: FLJ21930 fis, clone H	2.08 0.89	1.54 1.35
	330601 330605	U90916 X02419	Hs.82845 Hs.77274	plasminogen activator, urokinase	1.87	1.55
	330609	X04741	Hs.76118	ubiquitin carboxyl-terminal esterase L1	1.83	1.30
40	330817	X53587	Hs.85266	integrin, beta 4 reticulocalbin 2, EF-hand calcium bindin	1.54 1.39	1.15 1.19
40	330630 330644	X78669 Y07755	Hs.79088 Hs.38991	S100 calcium-binding protein A2	3.83	1.13
	330650	Z68228	Hs.2340	Junction plakoglobin	1.25	0.95
	330660	AA347868	Hs.139293	ESTs, Weakly similar to ALU7_HUMAN ALU S	15.50 1.00	29.07 1.00
45	330692 330707	AA017045 AA133891	Hs.6702 Hs.293690	ESTs ESTs	0.20	1.35
	330715	AA233707	Hs.11571	Homo sapiens cDNA FLJ11570 fis, clone HE	0.12	1.40
	330717	AA233926	Hs.52620	Integrin, beta 8 ESTs	6.62 1.40	5.42 1.65
	330722 330740	AA243560 AA297746	Hs.34382 Hs.22654	Homo sapiens voltage-gated sodium channe	0.27	2.04
50	330742	AA400979	Hs.25691	receptor (calcitonin) activity modifying	0.44	0.90
	330744	AA406142	Hs.12393 Hs.29643	dTDP-D-glucose 4,6-dehydratase Homo sapiens cDNA FLJ13103 fis, clone NT	0.71 1.66	3.23 1.52
	330751 330760	AA428286 AA448663	Hs.30469	ESTs	0.52	0.90
<i></i>	330763	AA450200	Hs.274337	hypothetical protein FLJ20666	0.37	0.97
55	330786	D60374 T48536	Hs.49136 Hs.105807	ESTs, Moderately similar to ALU7_HUMAN A ESTs	0.78 0.23	0.84 3.17
	330790 330814	AA015730	Hs.265398	ESTs, Weakly similar to transformation-r	0.37	2.07
	330827	AA040332	Hs.12744	ESTs	1.60	1.00
60	330844	AA063037	Hs.66803	ESTs endogenous retroviral protease	0.93 1.02	1.16 1.03
00	330901 330931	AA157818 F01443	Hs.267319 Hs.284256	hypothetical protein FLJ14033 similar to	0.24	0.88
	330952	H02855	Hs.29567	ESTs	0.08	1.31
	330961 330968	H10998 H16568	Hs.7164 Hs.23748	a disintegrin and metalloprotelnase doma ESTs	1.29 0.48	1.26 0.96
65	331014	H98597	Hs.30340	hypothetical protein KIAA1165	0.29	0.74
	331046	N66563	Hs.191358	ESTs	0.99	8.56
	331060 331099	N75081 R36671	Hs.157148 Hs.83937	Homo saplens cDNA FLJ11883 fts, clone HE hypothetical protein	1.24 0.75	1.00 1.03
	331108	R41408	Hs.21983	ESTs	1.00	2.75
70	331131	R54797		gb:yg87b07.s1 Soares infant brain 1NiB H	6.04	10.68
	331135 331170	R61398 T23461	Hs.4197 Hs.159293	ESTs ESTs	0.80 2.63	0.96 4.29
	331180	T32446	Hs.6640	Human DNA sequence from PAC 75N13 on chr	1.78	2.71
75	331183	T40769	Hs.8469	ESTs	1.00	3.01
75	331203 331271	T82310 AA059347	Hs.82226	(NONE) glycoprotein (transmembrane) nmb	1.70 1.20	3.80 3.19
	331271	AA252079	Hs.63931	dachshund (Drosophila) homolog	0.31	1.30
	331327	AA281076	Hs.109221	ESTs	2.09	2.41
80	331341	AA303125	Hs.23240 Hs.46901	Homo saplens cDNA FLJ13496 fis, clone PL KIAA1462 protein	0.72 0.09	2.43 0.91
30	331359 331363	AA416979 AA421562	Hs.91011	anterior gradient 2 (Xenepus laevis) hom	1.02	0.87
	331378	AA448881	Hs.49282	hypothetical protein FiJ11088	1.03	1.23
	331384	AA456001 AA505135	Hs.93847 Hs.44037	NADPH oxidase 4 ESTs	1.40 1.80	1.00 3.93
85	331402 331422	F10802	Hs.163628	ESTs, Moderately similar to ALU7_HUMAN	1.65	1.89
-	-					

	WO 02/086443							
	331490	N32912	Hs.26813	CDA14	2.48	1.73		
	331531	N51343		gb:yz15g04.s1 Soares_multiple_scierosis_ gb:od74f04.s1 NCI_CGAP_Ov2 Homo sapiens	0.98 3.80	1.68 5.75		
	331547 331578	N54811 N67960	Hs.249989	ESTs	0.11	0.67		
5	331589	N71027	Hs.152618	ESTs	1.09	1.38		
-	331608	N89861	Hs.112110	PTD007 protein	0.93	0.76		
	331614	N92293	Hs.240272	EST	0.17	1.34		
	331668	W69707	Hs.58030	EST	2.24 1.00	3.82 1.24		
10	331671 331676	W72033 W79834	Hs.194695 Hs.58559	ras homolog gene family, member l ESTs, Wealdy similar to rhotekin [M.musc	0.08	1.07		
10	331681	W85712	Hs.119571	collagen, type III, alpha 1 (Ehlers-Dani	8.72	4.27		
	331692	W93592	Hs.152213	wingless-type MMTV integration site fami	0.94	0.54		
	331717	AA190888	Hs.153881	Homo sapiens NY-REN-62 antigen mRNA, par	1.57	1.34		
15	331718	AA191404	Hs.104072	ESTs	6.80 1.10	11.77 1.00		
15	331811 331820	AA404500 AA405970	Hs.301570 Hs.97996	ESTs transcription termination factor, mitoc	0.73	0.59		
	331831	AA412031	Hs.97901	EST	2.77	4.08		
	331852	AA418988	Hs.98314	Homo saplens mRNA; cDNA DKFZp586L0120 (f	0.23	0.93		
20	331943	AA453418	Hs.21275	hypothetical protein FLJ11011	0.36	1.88		
20	331969	AA460702	Hs.82772	collagen, type XI, alpha 1	1.00 3.04	1.00 3.87		
	331990 332002	AA478102 AA482009	Hs.139631 Hs.105104	ESTs ESTs	1.19	0.78		
	332027	AA489671	Hs.65641	hypothetical protein FLJ20073	1.27	1.03		
	332029	AA489697	Hs.145053	ESTs	0.30	1.62		
25	332033	AA489840	Hs.251014	EST	2.30	3.70		
	332048	AA496019	Hs.201591	ESTs KIAA1211 protein	0.17 1.35	0.52 1.23		
	332071 332074	AA598594 AA599012	Hs.205293	gb:ae41e11.s1 Gessler Wilms tumor Homo s	0.19	2.00		
	332083	AA600200	Hs.155546	KIAA1080 protein; Golgi-associated, gamm	0.31	1.18		
30	332085	AA600353	Hs.173933	nuclear factor I/A	0.30	1.50		
	332125	AA609861	Hs.312447	ESTs	0.22	0.62		
	332177	F10812	Hs.101433	ESTs	8.21 2.27	18.03 1.57		
	332180 332185	H03348 H10356	Hs.7327 Hs.101689	claudin 1 ESTs	0.09	1.18		
35	332203	H49388	Hs.317769	EST	8.05	5.02		
	332232	N48891	Hs.101915	Stargardt disease 3 (autosomal dominant)	0.78	0.85		
	332240	N54803	Hs.324267	ESTs, Weakly similar to putative p150 [0.96 2.40	1.23 3.74		
	332261 332275	N70294 R08838	Hs.269137 Hs.26530	ESTs serum deprivation response (phosphatidy)	0.27	0.75		
40	332280	R38100	Hs.146381	RNA binding motif protein, X chromosome	0.39	1.88		
	332299	R69250	Hs.21201	nectin 3; DKFZP566B0846 protein	5.24	12.76		
	332304	R74041	Hs.101539	ESTs	1.44	3.18		
	332314	T25862	Hs.101774	hypothetical protein FLJ23045	0.68 1.71	1.32 0.88		
45	332384 332434	M11433 N75542	Hs.101850 Hs.289068	retinol-binding protein 1, cellular Homo sapiens cDNA FLJ11918 fis, clone HE	0.43	0.86		
73	332445	T63781	Hs.11112	ESTs	0.68	1.00		
	332453	L00205	Hs.111758	keratin 6A	31.54	1.00		
	332458	M33493	Hs.250700	tryptase beta 1	0.51	1.00		
50	332504	AA053917 M17252	Hs.15108 Hs.278430	chromosome 14 open reading frame 1 cytochrome P450, subfamily XXIA (steroid	0.79 0.98	1.24 1.70		
50	332525 332530	M31682	Hs.1735	Inhibin, beta B (activin AB beta polypep	0.88	0.66		
	332535	N20284	Hs.19280	cysteine-rich motor neuron 1	0.22	1.46		
	332539	AA412528	Hs.20183	ESTs, Weakly similar to AF164793 1 prote	0.93	1.49		
55	332559	M13955	Hs.166189	cytokeratin 2	0.35 1.00	1.13 1.00		
33	332563 332565	N92924 AA234896	Hs.274407 Hs.25272	protease, serine, 16 (thymus) E1A binding protein p300	0.36	1.05		
	332594	AA279313	Hs.3239	methyl CpG binding protein 2 (Rett syndr	0.53	0.59		
	332634	S38953	Hs.283750	tenascin XA	0.38	1.16		
60	332638	AA283034	Hs.50640	JAK binding protein	1.00	1.70 1.16		
60	332640 332654	AA417152 AA001296	Hs.5101 Hs.288217	protein regulator of cytokinesis 1 hypothetical protein MGC2941	6.15 1.50	2.73		
	332665	AA223335	Hs.63788	propionyl Coenzyme A carboxylase, beta p	1.20	0.91		
	332692	AA496035	Hs.247926	gap junction protein, alpha 5, 40kD (con	0.17	1.12		
CE	332716	L00058	Hs.79070	v-myc avian myelocytomatosis viral oncog	1.00	1.44 1.81		
65	332736	L13773 X93921	Hs.114765 Hs.296938	myeloid/lymphold or mixed-lineage leukem dual specificity phosphatase 7	1.00 0.53	0.78		
	332758 332781	AA233258	Hs.247112	hypothetical protein FLJ 10902	1.44	1.56		
	332792	74		.,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	1.70	1.19		
~^	332816				1.85	2.47		
70	332858				1.04 3.48	1.57 8.04		
	332906 332911				1.00	1.00		
	332912				1.06	4.40		
	332922				1.00	1.00		
75	332956				0.42	0.88		
	332959			•	1.96 0.56	6.34 0.99		
	332982 332984				0.30	0.78		
	332998				1.47	2.01		
80	333058				0.47	1.38		
	333097				2.14 2.76	3.19 3.70		
	333121				1.92	1.21		
	333122 333123				1.85	1.39		
85	333138				0.47	0.52		

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	333139		1.88
	333140		0.21 1.51
	333221 333260		0.75
5	333380		6.68
_	333387		4.56
	333512		5.05 2.28
	333524 333585		2.31
10	333603		2.23
	333604		2.51
	333618		0.52 1.44
	333627 333628		1.90
15	333650		1.85
	333678		1.85
	333750		2.18 1.99
	333763 333767		1.02
20	333768		1.78
	333769		2.15 1.46
	333772		1.46 1.00
	333777 333846		1.00 2.99 0.47
25	333884		0.47
	333887		0.50 0.43
	333891 333892	•	0.51
	333904		0.51 0.26
30	333906		0.55
	333948		1.70 0.37
	333954 333966		8.10
	333968		0.63
35	334061		4.24 1.30
	334094 334113		4.55
	334161		0.82
40	334183		0.47
40	334187		1.36 . 0.69
	334219 334222		1.88
	334223		4.72
15	334239		0.79 0.45
45	334255 334333		1.00
	334378		3.98
	334382		1.50 3.59
50	334492 334562		5.94
50	334588		8.14
	334616		1.55
	334633		5.16 0.59
55	334648 334787		3.70
	334866		8.13
	334891		0.32 1.00
	334933 334934		4.01
60	334945		1.04
	334967		0.29
	334990		1.50 5.88
	335015 335093		0.55
65	335120		4.31
	335125		0.38 1.24
	335179	•	0.46
	335188 335211		1.61
70	335288		0.73
	335289		0.20 2.18
	335361 335379		0.50
	335414		3.64
75	335416		2.93
	335496		0.96 1.71
	335497 335548		1.15
•	335551 .		3.22
80	335558		3.42 5.50
	335586 335619		2.99 ·
	335620		3,80
85	335621		0.28 0.46
03	335682	,	0.40

	WO 02/086	5443		
	335686		2.55	3.81
	335755		2.24	1.07
	335784		0.20	0.97
_	335814		1.13	1.48
5	335815		2.45	3.51
	335823		1.00	4.16
	335835		0.49	1.70
	335851		1.66	1.39
4.0	335868		2.98	6.43
10	335896		0.98	0.99
	335936		12.10	21.93
	335948		1.00	1.64
	335983		1.00	4.21 1.17
1.5	335995		0.37	
15	336021		1.04	0.84
	336034	•	11.40	23.54
	336038		1.19	1.21
	336066		0.54	1.63
20	336107		0.95	0.70
20	336205		3.13	6.29 10.10
	336275		3.20 2.34	3.09
	336292		1.00	1.00
	336331		0.65	0.79
25	336419		2.33	2.16
23	336632		2.55	2 23
	336633		2.19	2.23 2.03
	336634		2.69	2.48
	336635		2.13	1.83
30	336636		2.43	2.24
50	336637 336638		2.31	2.03
	336659		0.60	1.31
	336675		0.31	1.18
	336684		1.50	1.14
35	336694		4.74	7.10
55	336716		4.43	6.37
	336721		2.20	0.74
	336798		1.64	2.14
	336900		6.14	12.73
40	336948		1.00	1.00
	337028		1.30	2.09
	337043		4.01	11.53
	337046		1.67	1.84
	337054		2.78	7.35
45	337128		7.20	16.14
	337162		3.45	5.34
	337183		5.72	11.41
	337184		3.72	5.90
	337192		1.27	1.06
50	337194		1.88	1.68
	337229		0.22 1.00	1.03
	337268		1.00	3.31
	337299		3.23	5.14
	337325		2.76	3.72
55	337389		5.80	10.42
	337493		2.06	6.30
	337497		7.88	20.29
	337500		3.80	4.48 2.31
60	337549		1.66 1.27	8.54
60	337603		5.76	7.16
	337605		0.73	0.97
	337671		1.54	0.92
	337755		5.07	9.73
65	337786		6.18	12.87
05	337809		3.78	12.97
	337862		2.66	8.16
	337871 337958		0.26	1.34
	338008		1.48	1.12
70	338033		2.38	14.59
, 0	338083		0.65	2.16
	338110		1.00	1.61
	338112		5.86	8.25
	338145		1.70	1.97
75	338148		8.07	18.19
	338158		1.30	4.55
	338161		2.58	3.57
	338179		1.00	1.00
	338182		3.32	4.63
80	338189		1.00	3.34
	338197		0.99	1.69
	338199		4.58	7.62
	338215		6.01	15.85
	338279		0.53	0.95
85	338316		20.58	38.66

	338322	3.23	7.39
	338357	4.10	11.39
	338359	10.12	21.59
	338366	0.69	1.02
5	338374	0.40	1.18
	338414	0,47	1.06
	338418	6.12	13.86
	338469	3.09	5.11
	338501	6.28	10.32
10	338506	6.97	12.41
10	338523	3.10	5.84
	338549	1.70	2.70
	338561	0.79	0.81
	338662	1.72	1.46
15	338671	0.17	0.91
10	338676	2.10	15.86
	338726	1.20	1.09
	338779	0,12	0.57
	338804	0.99	1.67
20	338836	1.00	1.00
	338871	4.30	9.81
	338872	5.02	12.81
	338879	. 0.23	1.12
	338937	6.55	12.26
25	338966	1.76	5.42
	338993	1.00	2.40
	339047	5.26	10.81
	339100	5.10	6.88
	339114	1.00	1.70
30	339121	1.00	3.75
	339170	10.36	19.67
	339229	4.08	13.48
	339264	2.64	3.83
	339293	1.73	1.94
35			

TABLE 8B shows the accession numbers for those Pkeys in Table 8A lacking unigenelD's. For each probeset we have listed the gene cluster number from which the oligonucleotides were designed. Gene clusters were compiled using sequences derived from Genbank ESTs and mRNAs. These sequences were clustered based on sequence similarity using Clustering and Alignment Tools (DoubleTwist, Oakland California). The Genbank accession numbers for sequences comprising each cluster are listed in the "Accession" column.

CAT number: Gene cluster number Accession: Genbank accession numbers 45 CAT number Accessions Pkey AW340926 AA249063 N86075 Al341937 AW003063 U34725 AA904742 X57414 X57415 X13075 X13076 R93901 AF075073 R93902 322044 187363_1 322060 44320_1 50 321430 42705_1 43034_1 46779_1 321467 322125 R93901 AF075073 R93902 H69434 AF085958 H69846 H52567 H52567 AF085970 H52164 H56535 AF085980 H56712 H92891 AF085982 H92777 H84849 H84252 H84260 H86664 H85320 H95531 H95521 H84529 AA070412 AA102346 AA081885 H22544 H46842 AI204929 W69304 AF086283 W69200 AA625149 AA313030 AA313052 H97463 AA655089 AA135130 AA484059 AA102419 AW877765 W79150 AF086419 322166 46861_1 46873_1 46882_1 46885_1 322173 55 322178 322179 321577 1615102_1 1615333_1 111953_1 627492_1 47271_1 321587 313723 320997 60 322278 218439_1 129439_1 47422_1 321687 313883 322320 W79150 AF086419 W1913 AP00419 A1668646 A1734214 W17348 AW979268 AA878419 AA431342 AA431628 A1308300 A1308296 65 322339 814584_1 314648 300201 293660_1 682222_1 306897 25196_-2 AI093967 AL120701 AL135041 AL121524 AF147359 T58511 T58560 W88919 W89125 323155 979809_1 70 322527 322585 38927_1 473768_2 473768_2 1574395_1 82296_1 85042_1 380580_1 Z42308 H23514 300362 242305 H23514
A005129 AA679084 AA694399
AA011522 AA702841 AA011691 AA330797
A1239464 A1239473 AA625812 A1208703
AF074666 A1110759 AF090902
A1903735 AA491283 A1694953 AW976903 AA761362
A1347274 AW844024 322635 322664 315454 75 322687 37372_1 314852 327472_1 307783 697809_1 AJ347274 AW844024 AA381722 AA381829 AW983906 AW963902 AA381242 AA488472 W27363 AA317053 BE082699 AW967036 BE079872 AW970512 AA280251 Al652287 BE466438 Al650725 AA551854 AA281574 AW571481 AA678177 AA677034 324072 269032_1 80 300527 221345_1 323505 315791 196389_1 403558_1 324303 233842_1 AL118754 AA333202 H38001 316519 442885_1 AAB47835 AA768378 85 300926 333127 1 AA504860 AA504911

Unique Eos probeset identifier number

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	324580	328264_1	AA492588 AA492498 AA492571 .
	301882	275087_1	T78054 T79888 AA398185
	324804	398093_1	A1692552 A1393343 A1800510 A1377711 F24263 AA661876
	324889	1515978_1	D31010 D30991 D31168 D31166 D31465
5	302697	43219 1	AJ001409 AJ001410
•	302711	45419_1	L08442 D51348
	302742	458_39	L12061
	318499	364430_1	T25451 AA585296 AA585305
	310624	34624_4	U88896 U88898 AA916056 T03285 Al341594 Al359534 Al634031 U88897
10	302847	458_105	X98941 X98942 X98943 X98953 X98949
	304122	772715	H28966
	303598	270283_1	AA382814 AA402411 AA412355
	311409	837264_1	A1698839 A1909260 A1909259
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	319425	1689571_1	T82930 R02424 T85145
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	318865	1535937_1	H10818 F07831 Z43072
	312220	1671607_1	N74613 T98756 T98589
	319546	243305_1	R09692 R09414 AA346353
25	312389	902067_1	AIBG3140 W80703 R43474
25	319611	1566863_1	H14957 R56522 R11908 BE080180 AW827313 AW231970 AA995028 AA428584 AW872716 AW892508 AW854593 AA578441 AW975234 AA864937 AA984131
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	321102 321158	410938_1	H79670 H47798 AA700289
	321199	212379_1	N34524 AA305071 AW954803 AA502335 Al433430 Al203597 AW026670 AW265323 AW850787 AA317554 AW993643 AW835572
	321133	212373_1	AW385512 Al334966 W32951 H62656 H53902 R88904 AW835732
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	306458	AA978186	
	306510	AA988546	
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	306572	AA995686	
45	306582	AA996248	
	306656	AI004024	
	306686	AI015615	
	306751	AI032589	
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	308106	Al476803	
	308154	A1500600	
	306956	Al125111	
55	306958	Al125152	
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	308216	A1557135	·
	308219	Al557246 Al718299	
	308588 308599	Al719893	
60	308643	AI745040	
00	308673	A1760864	
	308697	Al767143	
	308778	AI811109	
	308808	Al818289	\cdot
65	308875	AI832332	
	308886	AI833240	
	308898	AI858845	
	308966	AI870704	•
	308979	Al873111	
70	303011	41689_1	AF090405 AF090407 AF090406
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	305016	AA626876	
	305034	AA630128	
	305072	AA641012	
75	305148	AA654070	
	305190	AA665955	
	303978	AW513315	
	303990	AW515465	·
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80	303999	AW516611	
	305235	AA670480	
	305312	AA700201	
	305413	AA724659	
85	305447	AA737856	AF068654 AF068656 AF068655
03	321244	29327_1	AL000034 AL000036 AL000039

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	305614 305637	AA782866 AA806124		•
	305639	AA806138		
_	305650	AA807709		
5	305690 305728	AA813477 AA828209		
	305759	AA835353		
	305792	AA845256		
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10	307091 307181	Al167439 Al189251		
	305901	AA872968		
	305910	AA875981		
15	307415 307426	Al242118 Al243364		
	307517	Al275055		
	307551	Al281556		
	307561 307608	A1282207 A1290295		
20	307691	Al318285		
	307730	Al336092		
	307760 307764	Al342387 Al342731		
	307796	Al350556		
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	309051 307807	Al911975 Al351799		
	307808	Al351826		
20	307820	Al355761		
30	307852 309122	Al365541 Al928178		
	309164	A)937761		
	309177	Al951118		
35	307902 309299	AJ380462 AW003478		
55	309303	AW004823		
	309476	AW129368		
	309532 309747	AW151119 AW264889		
40	309769	AW272346		
	309799	AW276964		
	309866	AW299916	HEEDOO A	A 4 0 C 0 C D
	302679 309923	311853_1 AW340684	H65022 A	A100009
45	309928	AW341418		
	309931	AW341683		
	309933 302705	AW341936 31765_1	U09060 U	09061
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50	304006	AW517947		
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	304028	T03266		
55	304046	T54803 T61521		
33	304061 304063	T62536		
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	304114 304155	R78946 H68696		
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	304234	W81608		
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	304456	AA411240		
65	304521	AA464716		
	304526 304607	AA476427 AA513322	•	
	304735	AA576453		
70	304760	AA580401		
70	306015 306063	AA897116 AA906316		
	306065	AA906725		
	306104	AA910956		
75	306109	AA911861		
13	306242 306288	AA932805 AA936900		
	306396	AA970223		
	330568	NOT_FOUND		U56244
80	330599 331131	1532312 genbank_R541	U90437 797	R54797
	331203	NOT_FOUND	_entrez	T82310
	331531	genbank_N51		N51343
	331547 332074	467396_1 a genbank_AA5	AA828597 99012	N54811 AA59901
85		Roundlin Add.		14-00001

TABLE 8C shows the genomic position for those Pkeys in Table 8A lacking unigene ID's and accession numbers. For each predicted exon, we have listed the genomic sequence source used for prediction. Nucleotide locations of each predicted exon are also listed.

Unique number corresponding to an Eos probeset
Sequence source. The 7 digit numbers in this column are Genbank Identifier (GI) numbers. "Dunham I. et al." refers to the publication entitled "The DNA sequence of human chromosome 22." Dunham I. et al., Nature (1999) 402:489-495.
Indicates DNA strand from which exons were predicted.
Indicates nucleotide positions of predicted exons. 5 Pkey: Ref:

Strand: Nt_position:

10	псроявон.	muicatos n	ucisoude poa	nuons of produ	cied exons.
	Pkey	Ref	Strand	Nt_position	
	000700	5 1 1 1		CI.	70004 70700
	332792 332816	Dunham, I. et.: Dunham, I. et.:		Plus Plus	73381-73768 359844-360030
15	332906	Dunham, I. et.		Plus	1923101-1923205
	332911	Dunham, I. et.		Plus	1961767-1961858
	332912	Dunham, I. et.		Plus	1962120-1962246
	332922	Dunham, I. et.:		Plus	2009620-2009738
20	332956	Dunham, I. et.:		Plus	2510528-2510658
20	332959 333138	Dunham, I. et.: Dunham, I. et.:		Plus Plus	2518145-2518213 3369205-3369323
	333139	Dunham, i. et.		Pius	3369495-3369571
	333221	Dunham, I. et.		Plus	3978070-3978187
0.5	333380	Dunham, I. et.a		Plus	4904775-4904846
25	333387	Dunham, I. et.a		Plus	4910935-4910997
	333512	Dunham, I. et.a		Plus	5560510-5560564
	333524 333585	Dunham, I. et.a Dunham, I. et.a		Plus .Plus	5612620-5612780 6234778-6234894
	333618	Dunham, I. et.		Plus	6562391-6562566
30	333627	Dunham, I. et.a		Plus	6620584-6620903
	333628	Dunham, I. et.a	al.	Plus	6629004-6629233
	333650	Dunham, I. et.a		Plus	6796852-6797128
	333678 333750	Dunham, I. et.a		Plus	7068223-7068288
35	333763	Dunham, I. et.a Dunham, I. et.a		Plus Plus	7608165-7608234 7692491-7692630
	333767	Ounham, I. et.a		Plus	7694407-7694623
	333768	Dunham, I. et.a		Plus	7695440-7695697
	333769	Dunham, i. et.a		Plus	7696625-7696707
40 .	333772	Dunham, I. et.a		Plus Plus	7706773-7706902
40 .	333777 333846	Dunham, I. et.a Dunham, I. et.a		Pius Plus	7746805-7746916 8008623-8008757
	333884	Dunham, I. et.a		Plus	8153960-8154161
	333887	Dunham, I. et.a		Plus	8154882-8155025
15	333891	Dunham, I. et.a		Pius	8156437-8156709
45	333892	Dunham, I. et.a Dunham, I. et.a		Plus Plus	8156825-8157001 8583497-8583627
	333948 333954	Dunham, I. et.a		Pius	6563186-6563335
	333966	Dunham, I. et.a		Plus	8655643-8655826
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50	334061	Dunham, I. et.a		Pius	9686941-9687077
	334094 334113	Dunham, I. et.a Dunham, I. et.a		Plus Plus	9889953-9890105 10282459-10282597
	334161	Dunham, I. et.a		Plus	10599033-10599180
	334219	Dunham, I. et.a	zi.	Plus	12716160-12716384
55	334239	Dunham, I. et.a		Plus	13056569-13056693
	334333 334378	Dunham, I. et.a		Plus Plus	13603544-13603657 13907239-13907370
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	335379	Dunham, I. et.a		Plus	22899306-22899420
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	335983	Dunham, I. et.a		Plus	27938968-27939070
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	336635	Dunham, I. et.al.	Plus	987908-988364
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• •	336637	Dunham, I. et.al.	Plus	989276-990813
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20	337183 337184	Dunham, I. et.al. Dunham, I. et.al.	Plus Plus	23943606-23943696 23973949-23974016
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	338279	Dunham, I. et.al.	Plus Plus	16168944-16169091 17089711-17089988
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ou	334223 334255	Dunham, I. et.al. Dunham, I. et.al.	Minus Minus	13200776-13200692
	334492	Dunham, I. et.al.	Minus	14478333-14478172
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OJ	334933	Dunham, I. et.al.	Minus	20078117-20077991

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1.5	335835	Dunham, I.		Minus Minus	26393311-26393245
	335851	Dunham, I.		Minus	26604863-26604742
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	336066	Dunham, I.		Minus	29241080-29240842
	336205 336275	Dunham, I. Dunham, I.		Minus Minus	30477456-30477311 32086675-32086536
25	336292	Dunham, I.		Minus	32818035-32817927
	336331	Dunham, I.		Minus	33594527-33594371
	336419 336675	Dunham, I. Dunham, I.		Minus Minus	34052568-34052445 2020758-2020664
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	336798 337043	Dunham, i. Dunham, i.		<i>Mi</i> nus Minus	5888954-5888757 17407330-17407251
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	325235 329588	6381943 3962484	Minus Plus	162154-162 1169-1619	2264
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70	325328 325340	5866875 6017033	Plus Minus	86780-8689 166656-166	
	325373	5866920	Minus	1136686-11	
	325367	5866920	Minus	922881-922	
75	325389 325436	5866921 5866939	Plus Minus	239672-239 29778-2990	
	325498	5866967	Plus	173372-173	1930
	325471	6017034	Minus	289268-289	
	325557 325559	6056302 6249595	Plus Minus	50921-5105 118590-119	
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	325569	6249599	Plus	79927-8021	
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	325969 325971	5867153 5867153	Plus Plus	105841-106035			
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	330020 326163	6671887 5867168	Plus Minus	172397-172491 7831-8035			
	326274	5867171	Minus	410289-410404			
25	326025	5867176	Plus	70854-70915			
25	326046 326099	5867182 5867186	Minus Minus	62668-62825 661381-661510			
	326108	5867187	Minus	23784-23903			
	326165	5867208	Minus	62787-62929			
30	326189 326204	5867212 5867218	Plus Minus	69288-69413 148088-148200			
50	326230	5867230	Minus	301868-301972			
	330052	4567182	Plus	352560-352963			
	330036 326360	6042048 5867293	Plus Plus	117120-117216 13627-13844			
35	326589	5867320	Plus	22760-22919			
	326393	5867341	Plus Minus	41702-41841 8818-8949			
	326505 326515	5867435 5867439	Plus	36683-36809			
40	326592	6138928	Plus	23689-23828			
40	330107 330106	6015249 6015249	Minus Minus	100091-100282 99443-99778			
	330100	6015253	Plus	21166-21301			
	330093	6015278	Plus	1043-1199			
45	330088 330085	6015293 6015302	Plus Minus	37517-37638 59613-59770			
	330120	6671864	Minus	127553-127656			
	330123 326742	6671869 5867611	Minus Minus	35311-35406 95187-95248			
	326605	5867637	Plus	24656-24749			
50	326818	6117831	Minus	15199-15309			
	326720 326770	6552456 6598307	Plus Minus	84525-84677 513603-513668			
	326692	6682502	Plus	117697-117899			
55	326693	6682502	Minus	335002-335095			
55	326983 326991	5867657 5867660	Minus Plus	16023-16581 18147-18339			
	326936	6004446	Minus	10217-10357			
	326964 327040	6469836 6531965	Plus Plus	75340-75456 783670-783817			
60	327053	6531965	Plus	2247267-2247437			
	327075	6531965	Plus	4041318-4041431			
	327085 327036	6531965 6531965	Plus Plus	4734947-4735069 319951-320040			
	327130	6531976	Pius	20247-22343			
65	327156	5866841	Minus	2462-2620			
	327288 327332	5867481 5867516	Plus Minus	48583-48773 56361-56532			
	327220	5867525	Minus	65701-65781			
70	327224	5867534	Plus	188468-188544			
70	327321 327361	6249562 6552412	Minus Minus	99745-99836 61013-62130			
	327396	5867743	Plus	8702-8820			
	327414	5867750	Plus	102461-102586			
75	327442 327467	5867759 5867772	Plus Plus	111483-111618 88030-88151			
	327473	5867775	Plus	75101-75181			
	327483 327377	5867783 5867793	Plus Minus	181573-181662 37610-37676			
0.0	327562	5867804	Minus	343989-344474			
80	327568	5867811	Minus	46152-46287			
	327606 327611	6004463 5867868	Plus Minus	200262-200495 175063-175392			
	327642	5867891	Minus	2513-2743			
85	327654	5867910	Minus	97564-97710			
3)	327734	5867940	Minus	31003-31583			

	,,,	, ,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,		
	327775	5867964	Minus	130791-130871
	327796	5867982	Plus	85267-85405
	327840	6249578	Minus	73065-73206
_	330208	6013599	Plus	66517-66931
5	330263	6671884	Minus	101503-101634
	328004	5867993	Minus	157407-157887
	328101	5868020	Plus	289920-290014
	328100	5868020	Minus	263545-263635
	328113	5868024	Minus	80378-80491
10	328157	5868064	Plus	73326-73615
	328196	5868080	Minus	16551-16729
	328197	5868081	Minus	42133-42438
	327940	5868197	Minus	95240-95428
	327984	5868216	Plus	66611-66677
15	328021	5902482	Plus	713478-714590
	328068	6117819	Plus	253903-254022
	328264	6381912	Plus	55086-55404
	330300	2905862	Minus	3246-3302
	328608	5868222	Minus	87770-87953
20	328600	5868229	Minus	38889-40010
	328616	5868239	Plus	293920-294224
	328623	5868246	Minus	120020-120126
	328632	5868247	Plus	76734-76853
~ -	328666	5868254	Minus	778-901
25	328698	5868264	Minus	625555-625633
	328700	5868264	Plus	764089-764203
	328708	5868271	Minus	68114-68854
	328735	5868289	Pius	89389-89455
20	328743	5868289	Plus	274638-274726
30	328806	5868324	Plus	29408-29684
	328299	5868366	Minus	149708-149889
	328342	5868383	Plus	59955-60094
	328365	5868387	Minus	270724-270798
35	328369	5868388	Plus	75371-75583
33	328381	5868392	Plus	662758-662848 217275-217336
	328451	5868425 5868449	Minus Minus	8987-9180
	328481	5868464	Pius	59098-69481
	328500 328530	5868482	Plus	334973-335406
40	328664	6004473	Plus	1193739-1193866
70	328861	6381928	Minus	108317-108403
	328908	5868493	Pius	117002-117059
	328933	5868500	Plus	771755-771889
	328934	5868500	Plus	846342-846448
45	328949	6456765	Minus	43552-43619
	330313	6042030	Minus	33642-33775
	329005	5868542	Plus	85470-85673
	330366	2944106	Plus	151837-151914
	330372	6580495	Minus	317461-317688
50	329033	5868561	Minus	5390-5479
	329037	5868562	Minus	32466-32562
	329067	5868591	Minus	146417-147652
	329134	5868679	Plus	29959-30018
	329157	5868687	Minus	145940-146155
55	329178	5868704	Plus	179177-179463
	329192	5868716	Plus	166936-167020
	329194	5868716	Minus	304450-304559
	329204	5868720	Minus	3050-3190
	329224	5868728	Plus	27422-27664
60	329228	5868728	Minus	50118-50287
	329288	5868771	Plus	25554-26299
	329337	5868806	Minus	467155-467222
	329011	6682532	Plus	48658-48741

TABLE 9A: Potential Therapeutic, Diagnostic and Prognostic targets for Therapy of Lung Cancer

Table 9A shows about 1312 genes up-regulated in lung tumors (including squamous cell carcinomas, adenocarcinomas, small cell carcinomas, granulomatous and carcinold tumors) relative to normal body tissues. These genes were selected from about 59680 probesets on the Eos/Affymetrix Hu03 Genechip array.

Table 9B show the accession numbers for those Pkey's lacking UnigenelD's for table 9A. For each probeset we have listed the gene cluster number from which the oligonucleotides were designed. Gene clusters were compiled using sequences derived from Genbank ESTs and mRNAs. These sequences were clustered based on sequence similarity using Clustering and Alignment Tools (DoubleTwist, Oakland California). The Genbank accession numbers for sequences comprising each cluster are listed in the "Accession" column.

Table 9C show the genomic positioning for those Pkey's lacking Unigene ID's and accession numbers in table 9A. For each predicted exon, we have listed the genomic sequence source used for prediction. Nucleotide locations of each predicted exon are also listed.

Exempla Accession number, consum accession in	15	Pkey: ExAccn:	•	Unique Eos probeset identifier number Exemplar Accession number, Genbank accession nur	nbe
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UnigeneID: Unigene number

5

10

Unigene Title: Unigene gene title

Average of lung tumors (including squamous cell carcinomas, adenocarcinomas, small cell carcinomas, granulomatous and carcinoid tumors) divided by the average of normal lung samples

Average of non-mailgnant lung disease samples (including bronchitis, emphysema, fibrosis, atelectasis, asthma) divided by the average of normal lung samples

20

R2:

	Pkey	ExAcon	UnigenelD	Unigene Title	R1	R2
	400195	LAWOII	Oligenero	NM_007057*:Homo sapiens ZW10 interactor	1.00	1.00
25	400205			NM_006265*:Homo sapiens RAD21 (S. pombe)	15.80	396.00
	400220			Eos Control	2.28	2.84
	400277			Eos Control	7.68	9.72
	400285			Eos Control	1.00	1.00
	400288	X06256	Hs.149609	integrin, alpha 5 (fibronectin receptor,	1.04	2.24
30	400289	X07820	Hs.2258	matrix metalloproteinase 10 (stromelysin	132.45	4.00
	400298	AA032279	Hs.61635	six transmembrane epithelial antigen of	43.86	74.00
	400301	X03635	Hs.1657	estrogen receptor 1	1.00	1.00
	400303	AA242758	Hs.79136	LIV-1 protein, estrogen regulated	1.75	1.65
~ -	400328	X87344	Hs.180062	transporter 2, ATP-binding cassette, sub	0.87	1.80
35	400419	AF084545		Target	156.55	253.00
	400512			NM_030878*:Homo sapiens cytochrome P450,	1.00	2.00
	400517	AF242388		lengsin	3.67	87.00
	400560			NM_030878*:Homo sapiens cytochrome P450,	1.00	1.00 45.00
40	400664			NM_002425:Homo sapiens matrix metallopro	20.26	1.07
40	400665			NM_002425:Homo sapiens matrix metallopro	1.36 3.26	3.22
	400666			NM_002425:Homo sapiens matrix metallopro	1.00	91.00
	400749			NM_003105*:Homo sapiens sortilin-related	7.63	24.00
	400763			Target Exon	1.00	1.00
45	401027 401093			Target Exon C12000586*:gl]6330167[db] BAA86477.1] (A	1.00	155.00
47	401093			Target Exon	1.00	86.00
	401212			C12000457*:gi[7512178]pir][T30337 polypr	1.00	400.00
	401411			ENSP00000247172*:HYPOTHETICAL 126.2 kDa	1.00	72.00
	401435			C14000397*:gi]7499898jpirl[T33295 hypoth	1.00	64.00
50	401464	AF039241		histone deacetylase 5	3.82	49.00
-	401714			ENSP00000241802°:CDNA FLJ11007 FIS, CLON	2.02	40.00
	401747			Homo saplens keratin 17 (KRT17)	128.43	68.00
	401760			Target Exon	1.74	35.00
	401780			NM_005557*:Homo sapiens keratin 16 (foca	26,47	10.50
55	401781			Target Exon	10.33	4.61
-	401785			NM_002275*:Homo sapiens keratin 15 (KRT1	4.13	2.70
	401797			Target Exon	1.44	2.10
	401961			NM_021626:Homo sapiens serine carboxypep	1.41	1.86
C O	401985	AF053004		class I cytokine receptor	1.00	177.00
60	401994			Target Exon	61.84	47.00
	402075			ENSP00000251056*:Plasma membrane calcium	1.00	1.00
	402260			NM_001436*:Homo sapiens fibrillarin (FBL	1.58 2.09	1.39 35.00
	402265			Target Exon	1.00	92.00
65	402297			Target Exon NM_030920*:Homo sapiens hypothetical pro	28.87	13.00
05	402408			C1000823*:gij10432400jembjCAC10290.1j (A	1.00	1.44
	402420 402674			Target Exon	7.44	243.00
	402802			NM_001397:Homo sapiens endothelin conver	1.00	70.00
	402994			NM_002463*:Homo sapiens myxovirus (influ	1.37	1.43
70	403137			NM_005381*:Homo sapiens nucleolin (NCL),	1.00	19.00
, 0	403306	NM_006825		transmembrane protein (63kD), endoplasmi	1.00	43.00
	403329			Target Exon	1.00	61.00
	403381			ENSP00000231844*:Ecotropic virus integra	1.00	119.00
_	403478			NM_022342:Homo sapiens kinesin protein 9	28.13	136.00
75	403485			C3001813*:gij12737279 ref XP_012163.1 k	20.23	76.00
	403627			Target Exon	6.30	29.33
	403715			Target Exon	1.30	35.00
	404044			ENSP00000237855*:DJ398G3.2 (NOVEL PROTE)	1.00	54.00
00	404076			NM_016020*:Homo sapiens CGI-75 protein (14.29	91.00
80	404101			C8000950:gi[423560 plr][A47318 RNA-bindi	1.00	1.00
	404140			NM_006510:Homo sapiens ret finger protei	1.42	1.44
	404165			ENSP00000244562:NRH dehydrogenase [quino	1.00 1.00	54.00 117.00
	404185			Target Exon	7.00 5.93	13.77
85	404210			NM_005936:Homo saptens myelold/lymphoid NM_021058*:Homo saptens H2B histone fami	1.00	1.00
05	404253			HINTOS 1000 'LIOURO SANIGUS LISTO HISTORIA (SILII)	1.00	1.00

	W	O 02/086	443			
	404287			C6001909:gij704441]dbjjBAA18909.1] (D298	29.71	42.00
	404298			C6001238*:gij121715 sp P26697 GTA3_CHICK	1.30	1.00
	404347			Target Exon	1.00	1.00
~	404440			NM_021048:Homo sapiens melanoma antigen,	1.00	15.00
5	404721			NM_005596*:Homo sapiens nuclear factor I	1.00	60.00
	404794	NM_000078		cholesteryl ester transfer protein, plas	1.07	1.38
	404854	_		Target Exon	1.61	2.01
	404877			NM_005365:Homo sapiens melanoma antigen,	1.00	1.00
	404927			Target Exon	1.00	1.00
10					1.00	1.00
10	404996			Target Exon		
	405449			CY000047*:gi[11427234]ref[XP_009399.1] z	1.00	1.00
	405568			NM_031413*:Homo sapiens cat eye syndrome	1.00	78.00
	405572			Target Exon	0.76	1.14
	405646			C12000200:gi 4557225 ref[NP_000005.1] al	1.01	1.28
15	405676	BE336714		cytochrome c-1	1.13	2.89
10	405770	DE0001 14		NM 002362:Homo saglens melanoma antigen,	45.52	37.00
					1.99	1.99
	405932			C15000305:gi 3806122 gb AAC69198.1 (AF0		
	406137			NM_000179*:Homo sapiens mutS (E. coli) h	2.77	2.38
•	406360			Target Exon	1.00	35.00
20	406399			NM_003122*:Homo sapiens serine protease	1.00	39.00
	406467			Target Exon	1.00	1.00
	406621	X57809	Hs.181125	immunoglobulin lambda locus	1.41	1.74
	406642	AJ245210		gb:Homo sapiens mRNA for immunoglobulin	2.16	3.91
			Hs.293441	immunoglobulin heavy constant mu	2.07	2.93
25	406663	U24683				
25	406671	AA129547	Hs.285754	met proto-oncogene (hepatocyte growth fa	15.00	51.00
	406673	M34996	Hs.198253	major histocompatibility complex, class	0.98	3.09
	406676	X58399	Hs.81221	Human L2-9 transcript of unrearranged im.	1.30	1.53
	406678	U77534		gb:Human clone 1A11 immunoglobulin varia	1.33	1.45
	406685	M18728		gb:Human nonspecific crossreacting antig	1.46	2.85
30	406687	M31126	Hs.272822	pregnancy specific beta-1-glycoprotein 9	8.61	8.50
50		M29540		carcinoembryonic antigen-related cell ad	226.37	350.00
	406690		Hs.220529			
	406698	X03068	Hs.73931	major histocompatibility complex, class	1.01	2.52
	406815	AA833930	Hs.288036	tRNA isopentenylpyrophosphate transferas	20.25	32.00
	406851	AA609784		major histocompatibility complex, class	0.75	1.91 ·
35	406964	M21305		gb:Human alpha satellite and satellite 3	38.15	1114.00
	406967	M24349		gb:Human parathyroid hormone-like protei	1.00	1.00
	406974	M57293		gb:Human parathyroid hormone-related pep	1.00	1.00
	407103	AA424881	Hs.256301	hypothetical protein MGC13170	1.77	1.10
					1.00	1.00
40	407128	R83312	Hs.237260	EST		
40	407137	T97307		gb:ye53h05.s1 Soares fetal liver spleen	142.70	135.00
	407168	R45175	Hs.117183	ESTs	2.16	18.00
	407239	AA076350	Hs.67846	leukocyte immunoglobulin-like receptor,	1.10	1.57
	407242	M18728		gb:Human nonspecific crossreacting antig	1.12	2.85
	407244	M10014	Hs.75431	fibrinogen, gamma polypeptide	3.24	15.38
45	407289	AA135159	Hs.203349	Homo saplens cDNA FLJ12149 fis, clone MA	3.53	3.68
73					19.74	73.00
	407300	AA102616	Hs.120769	gb:zn43e07.s1 Stratagene HeLa cell s3 93		
	407366	AF026942	Hs.271530	gb:Homo sapiens cig33 mRNA, partial sequ	0.06	8.25
	407378	AA299264	Hs.57776	ESTs, Moderately similar to I38022 hypot	1.00	26,00
	407430	AF169351		gb:Homo sapiens protein tyrosine phospha	1.00	25.00
50	407453	AJ132087		gb:Homo saplens mRNA for axonemal dynein	1.00	75.00
	407577	AW131324	Hs.246759	hypothetical protein MGC12538	1.00	1.00
	407634	AW016569	Hs.136414	UDP-GicNAc:betaGal beta-1,3-N-acetylgluc	111.20	228.00
	407710	AW022727	Hs.23616	ESTs	1.00	28.00
		AB037776			1.89	1.31
55	407720		Hs.38002	KIAA1355 protein	1.00	1.00
23	407746	AK001962		hypothetical protein FLJ11100		
	407756	AA116021	Hs.38260	ubiquitin specific protease 18	4.51	5.00
	407758	D50915	Hs.38365	KIAA0125 gene product	1.00	28.00
	407782	AA608956	Hs.112619	ESTs, Moderately similar to PURKINJE CEL	0.97	1.14
	407788	BE514982	Hs.38991	S100 calcium-binding protein A2	7.88	3.83
60	407790	AI027274	Hs.288941	Homo sapiens cDNA FLJ14866 fis, clone PL	3.63	42.00
	407811	AW190902	Hs.40098	cysteine knot superfamily 1, BMP antagon	89.96	109.00
	407839	AA045144	Hs.161566	ESTs	173.91	108.00
		R34008		desmocollin 2	111.30	70.00
	407944		Hs.239727			
65	408000	L11690	Hs.620	bullous pemphigoid antigen 1 (230/240kD)	151.17	8.00
65	408031	AA081395	Hs.42173	Homo sapiens cDNA FLJ10366 fis, clone NT	9.91	93.00
	408063	BE086548	Hs.42346	calcineurin-binding protein calsarcin-1	195.78	231.00
	408070	AW148852		gb:xf05d05.x1 NCI_CGAP_Bm35 Homo sapien	1.00	1.00
	408101	AW968504	Hs.123073	CDC2-related protein kinase 7	37.84	61.00
	408122	A1432652	Hs.42824	hypothetical protein FLJ10718	0.85	1.71
70	408212	AA297567	Hs.43728	hypothetical protein	5.88	7.91
, 0					4.27	9.98
	408243	Y00787	Hs.624	interleukin 8		
	408349	BE546947	Hs.44276	homeo box C10	3.79	3.46
	408353	BE439838	Hs.44298	mitochondrial ribosomal protein S17	1.88	1.65
a	408354	Al382803	Hs.159235	ESTs	1.00	73.00
75	408369	R38438	Hs.182575	solute carrier family 15 (H??? transport	1.41	16.50
	408380	AF123050	Hs.44532	diubiquitin	15.19	37.22
	408482	NM_000676	Hs.45743	adenosine A2b receptor	1.65	1.19
		AI541214	Hs.46320	Small proline-rich protein SPRK (human,	1.98	1.24
	408522					
0Λ	408536	AW381532	Hs.135188	ESTs	1.55	1.50
80	408545	AW235405	Hs.253690	ESTs	1.00	1.00
	408572	AA055611	Hs.226568	ESTs, Moderately similar to ALU4_HUMAN A	1.00	44.00
	408633	AW963372	Hs.46677	PRO2000 protein	107.16	56.00
	408660	AA525775		ESTs, Moderately similar to PC4259 ferri	1.00	1.00
	408761	AA057264	Hs.238936	ESTs, Weakly similar to (defline not ava	52.24	141.00
85	408771	AW732573	Hs.47584	potassium voltage-gated channel, delayed	3.05	109.00
				harmon compa dama anamon agintas		

WO 02/086443 Hs.47701 408783 AF192522 NPC1 (Niemann-Pick disease, type C1, gen 1.02 1.07 61.00 Hs.47860 neurotrophic tyrosine kinase, receptor, 41.19 408790 AW580227 Hs.48269 vaccinia related kinase 1 45.00 408805 H69912 Hs.256862 1.00 58.00 408841 AW438865 5 408873 AL046017 Hs.182278 calmodulin 2 (phosphorylase kinase, delt 1.00 89 00 serine/hreonine kinase 15 guarine nucleotide binding protein (G pr glycoprotein (transmembrane) nmb NM_004553:Homo saplens NADH dehydrogenas 1.00 BE296227 408908 Hs.250822 7.76 Hs.71642 1.00 1.00 408992 AA059325 Hs.344096 5.50 Al979168 408996 1.24 Hs.49767 1 44 409015 small inducible cytokine subfamily A (Cy Hypothetical protein, XP_051860 (KIAA119 5.32 10 409038 T97490 Hs.50002 4.2R 112.42 195.00 AB033025 Hs.50081 409041 17.00 409077 AA401369 Hs.190721 1.00 BE243834 Hs.50441 CGI-04 protein 2.02 1.93 409093 XAGE-1 protein SMC4 (structural maintenance of chromoso 80.44 40.00 409103 AF251237 Hs.112208 15 6.00 14.87 409142 AL136877 Hs.50758 Hs.50986 carbamoyl-phosphate synthetase 1, mitoch ESTs, Weakly similar to 2109260A B cell 1.00 1.00 AF154830 409187 Hs.271695 1.22 1.00 409228 AI654298 409234 1.00 Al879419 Hs.27206 **ESTs** 1.00 11.90 23.00 409268 AA625304 Hs.187579 **ESTs** 20 AA576953 Hs.22972 hypothetical protein FLJ13352 1.00 1.00 409269 NM 005982 Hs.54416 sine oculis homeobox (Drosophila) homolo 168.91 35.00 409361 1.00 409404 BE220053 Hs.129056 ESTs 1.00 Hs.54451 409420 Z15008 laminin, gamma 2 (nicein (100kD), kalini 79.74 96,00 splicing factor, arginine/serine-rich 5 2.10 R21945 Hs.346735 409430 25 409446 Al561173 Hs.67688 **ESTs** 1.00 4.00 409506 NM_006153 Hs.54589 NCK adaptor protein 1 3.97 28.00 gb:zm87b03.s1 Stratagene ovarian cancer 15.98 141.00 AA075382 AA401369 409522 ESTs Hs.190721 1.00 17.00 409582 W74001 serine (or cysteine) proteinase inhibito 292.12 79.00 Hs.55279 409632 30 brain-derived neurotrophic factor Homo sapiens brain tumor associated prot 409705 M37762 Hs.56023 1.00 82.00 1.00 409719 A176916D Hs.108681 1.00 thymosin, beta, identified in neuroblast 409731 AA125985 Hs.56145 Homo saplens mRNA; cDNA DKFZp586P2321 (f 20.75 51.00 409744 AW675258 Hs.56265 NM_001898 AW502152 409757 Hs.123114 22,46 15.80 35 gb:UI-HF-BR0p-air-f-11-0-UI.r1 NIH_MGC_5 1.00 1.00 409866 minichromosome maintenance deficient (S. AW247090 Hs.57101 409893 409902 Al337658 Hs.156351 25.92 50.00 409935 AW511413 Hs.278025 **ESTs** 2 63 2.11 4.01 inhibin, beta A (activin A, activin AB a 2.17 409956 AW103364 Hs.727 40 hyaluronan synthase 1 NM_001523 Hs.57697 2.07 409958 AB041036 Hs.57771 kallikrein 11 1.04 2.28 410001 gb:RC3-BT0319-120200-014-a09 BT0319 Homo 58.00 410032 BE085985 1.00 Hs.58009 1.00 34.00 KIAA0918 protein highly expressed in cancer, rich in leuc 410037 AB020725 1.00 410044 BE566742 Hs.58169 45 410048 W76467 Hs.58218 proline oxidase homolog 1.03 1 44 1.50 410076 T05387 Hs.7991 ESTS 1.12 410102 AW248508 Hs.279727 Homo sapiens cDNA FLJ14035 fis, clone HE 410153 BE311926 Hs.15830 hypothetical protein FLJ12691 1.00 1.00 hypothetical protein FLJ 10514 410166 AK001376 Hs.59346 1.00 1.00 50 410193 AJ132592 Hs.59757 zinc finger protein 281 hypoxia-inducible protein 2 42.01 51.00 AA381807 410274 Hs.61762 1.00 BE043077 Hs.278153 2.00 410309 410340 AW182833 Hs.112188 hypothetical protein FLJ13149 32.08 75.00 1.00 1.00 410348 AW182663 Hs 95469 EST₈ 55 410407 carbonic anhydrase IX Hs.63287 X66839 transmembrane protease, serine 4 hypothetical protein KIAA1335 410418 D31382 Hs.63325 4.30 2.03 AB037756 AW016824 Hs.45207 Hs.255527 18.00 410438 1.00 hypothetical protein MGC14128 410553 a disintegrin and metalloproteinase doma 1.41 410555 Hs.64311 W27235 60 410561 BE540255 Hs.6994 Homo sapiens cDNA: FLJ22044 fis, clone H 10.04 1.00 10.88 18.92 410681 AW246890 Hs.65425 calbindin 1, (28kD) Hs.165028 57.00 Al375672 **ESTs** 410781 411027 AF072099 Hs.67846 leukocyte immunogłobulin-like receptor, 1.62 3.78 adenylate cyclase activating polypeptide cell division cycle 2-like 1 (PITSLRE pr gb:QV3-BT0379-010300-105-g03 BT0379 Homo Homo sapiens cDNA FLJ14408 fis, clone HE 411074 X60435 Hs.68137 1.00 1.15 65 1.58 411089 AA456454 1.58 BE069199 411152 Hs.334605 1.82 1.45 411248 AA551538 411252 AB018549 Hs.69328 MD-2 protein 7.32 12.74 kinesin-like 6 (mitotic centromere-assoc 3.44 2.55 411263 BE297802 Hs.69360 70 411365 GM2 ganglioside activator protein Hs.289082 M76477 NRAS-related gene 411402 BE297855 1.00 46.00 Hs.69855 KIAA1077 protein U6 snRNA-associated Sm-like protein LSm7 11.35 411573 AB029000 Hs.70823 11.40 Hs.70830 1.08 1.90 AC005258 411579 411617 AA247994 Hs.90063 neurocalcin delta 2.57 75 AA059325 Hs.71642 guanine nucleotide binding protein (G pr 1.02 1.00 411732 411773 NM_006799 AF245505 Hs.72026 protease, serine, 21 (testisin) 1.34 2.19 411789 Hs.72157 Adlican Hs.103042 microtubule-associated protein 18 34.00 411800 N39342 23.34 v-myc avian myelocytomatosis viral oncog hypothetical protein FLJ10901 RAB6 interacting, kinesin-like (rabkines 411945 AL033527 Hs.92137 1.00 8.00 80 1.64 412115 AK001763 Hs.73239 2.07 118.48 92.00 412140 AA219591 Hs.73825 macrophage migration inhibitory factor (ESTs, Weakly similar to 155214 salivary 412276 BE262621 Hs.73798 1.49 1.98 Hs.22826 1.16 1.34 412464 T78141 hypothetical protein FLJ13346 84.00 AA766268 41.52 412530 Hs.266273 85 nuclear transcription factor Y, alpha

412537

AL031778

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	412659	AW753865	Hs.74376	olfactomedin related ER localized protei	14.65	47.00
	412719 412723	AW016610 AA648459	Hs.816 Hs.335951	ESTs hypothetical protein AF301222	382.46 54.90	128.00 1.00
	412811	H06382	NS.333331	ESTs	1.00	11.00
5	412817	AL037159	Hs.74619	proteasome (prosome, macropain) 26S subu	1.63	1.42
	412863	AA121673	Hs.59757 Hs.75258	zinc finger protein 281 H2A histone family, member Y	17.63 1.00	56.00 22.00
	412924 413004	BE018422 T35901	Hs.75117	interleukin enhancer binding factor 2, 4	2.19	2.05
4.0	413011	AW068115	Hs.821	biglycan	1.22	1.88
10	413048	M93221	Hs.75182	mannose receptor, C type 1	0.30 3.43	6.23 8.71
	413063 413129	AL035737 AF292100	Hs.75184 Hs.104613	chitinase 3-like 1 (cartilage glycoprote RP42 homolog	4.67	4.77
	413142	M81740	Hs.75212	omithine decarboxylase 1	1.92	2.59
15	413223	AI732182 ·	Hs.191866	ESTs	5.73 0.99	27.00 1.06
15	413248 413273	T64858 U75679	Hs.21433 Hs.75257	hypothetical protein DKFZp547J036 stem-loop (histone) binding protein	1.00	18.00
	413278	BE563085	Hs.833	interferon-stimulated protein, 15 kDa	1.10	1.09
	413281	AA861271	Hs.222024	transcription factor BMAL2	95.94 1.00	69.00 1.00
20	413364 413385	BE536218 M34455	Hs.137516 Hs.840	fidgetin-like 1 indoleamine-pyrrole 2,3 dioxygenase	0.95	2.09
	413409	Al638418	Hs.1440	DEAD/H (Asp-Glu-Ala-Asp/His) box polypep	1.00	1.00
	413453	AA129640	Hs.128065	ESTs	1.00 1.08	31.00 1.46
	413527 413554	BE250788 AA319146	Hs.179882 Hs.75426	hypothetical protein FLJ12443 secretogranin II (chromogranin C)	79.15	114.00
25	413573	AI733859	Hs.149089	ESTs	1.00	1.00
	413582	AW295647	Hs.71331	hypothetical protein MGC5350	8.80 1.00	10.00 1.00
	413597 413690	AW302885 BE157489	Hs.117183	ESTs gb:RC1-HT0375-120200-011-e06 HT0375 Homo	1.00	1.00
••	413691	AB023173	Hs.75478	ATPase, Class VI, type 11B	3.16	2.32
30	413719	BE439580	Hs.75498	small inducible cytokine subfamily A (Cy	2.88 144.10	9.52 108.00
	413753 413801	U17760 M62246	Hs.75517 Hs.35406	laminin, beta 3 (nicein (125kD), kalinin ESTs, Highly similar to unnamed protein	1.00	17.00
	413833	Z15005	Hs.75573	centromere protein E (312kD)	1.00	1.00
35	413882	AA132973	Hs.184492 Hs.54310	ESTs	64.24 1.00	148.00 67.00
33	413926 413943	AA133338 AW294416	Hs.144687	ESTs Homo sapiens cDNA FLJ12981 fis, done NT	43.42	42.00
	413995	BE048146	Hs.75671	syntaxin 1A (brain)	1.23	1.11
	414035	Y00630	Hs.75716 Hs.334485	serine (or cystelne) proteinase inhibito Homo sapiens cDNA FLJ14438 fis, clone HE	2.02 1.00	2.51 102.00
40	414142 414180	AW368397 A1863304	Hs.120905	Homo sapiens cDNA FLJ11448 fis, done HE	6.92	77.00
	414245	BE148072	Hs.75850	WAS protein family, member 1	1.00	1.00
	414275 414317	AW970254 BE263280	Hs.889 Hs.75888	Charot-Leyden crystal protein phosphogluconate dehydrogenase	1.00 1.52	59.00 1.73
	414334	AA824298	Hs.21331	hypothetical protein FLJ10036	1.78	1.72
45	414341	D80004	Hs.75909	KIAA0182 protein	33.90	151.00
	414368 414416	W70171 AW409985	Hs,75939 Hs,76084	uridine monophosphate kinase hypothetical protein MGC2721	171.60 2.32	97.00 1.85
	414430	Al346201	Hs.76118	ubiquitin carboxyl-terminal esterase L1	226.15	66.00
50	414570	Y00285	Hs.76473	insulin-like growth factor 2 receptor	1.64 1.87	1.98 72.00
30	414618 414675	Al204600 R79015	Hs.96978 Hs.296281	hypothetical protein MGC10764 interleukin enhancer binding factor 1	1.51	1.39
	414683	S78296	Hs.76888	hypothetical protein MGC12702	43.61	64.00
	414696 414711	AF002020 A1310440	Hs.76918 Hs.288735	Niemann-Pick disease, type C1 Homo sapiens cDNA FLJ13522 fis, clone PL	28.63 14.86	71.00 42.00
55	414718	H95348	Hs.107987	ESTs	1.00	5.00
	414732	AW410976	Hs.77152	minichromosome maintenance deficient (S.	1.64	1.44
	414747 414761	U30872 AU077228	Hs.77204 Hs.77256	centromere protein F (350/400kD, mitosin enhancer of zeste (Drosophila) homolog 2	65.01 130.35	74.00 121.00
	414774	X02419	Hs.77274	plasminogen activator, urokinase	2.24	2.19
60	414806	D14694	Hs.77329	phosphatidylserine synthase 1 transferrin receptor (p90, CD71)	1.63 1.97	1.53 2.60
	414809 414812	A1434699 X72755	Hs.77358 Hs.77367	monokine induced by gamma interferon	3.48	10.60
	414825	X06370	Hs.77432	epidermal growth factor receptor (avian	103.22	143.00
65	414839 414883	X63692 AA926960	Hs.77462	DNA (cytosine-5-)-methyltransferase 1 CDC28 protein kinase 1	1.80 14.29	1.69 10.06
05	414907	X90725	Hs.77597	polo (Drosophia)-like kinase	1.95	2.20
	414914	U49844	Hs.77613	ataxia telanglectasia and Rad3 related	3.00	2.90 1.21
	414945 414972	BE076358 BE263782	Hs.77667 Hs.77695	tymphocyte antigen 6 complex, locus E KIAA0008 gene product	1.02 1.00	1.00
70	415014	AW954064	Hs.24951	ESTs	1.42	2.84
	415091	AL044872	Hs.77910	3-hydroxy-3-methylglutaryl-Coenzyme A sy	1.00 34.72	30.00 107.00
	415138 415227	C18356 AW821113	Hs.295944 Hs.72402	tissue factor pathway inhibitor 2 ESTs	1.87	49.00
~~	415238	R37780	Hs.21422	ESTs	1.00	1.00
75	415263	AA948033	Hs.130853 Hs.6546	ESTs ESTs	1.00 1.00	1.00 1.00
	415295 415339	R41450 NM_015156	Hs.78398	KIAA0071 protein	51.18	166.00
	415669	NM_005025	Hs.78589	serine (or cysteine) proteinase inhibito	30.84	63.00
80	415674	BE394784 AA649850	Hs.78596 Hs.278558	proteasome (prosome, macropain) subunit, ESTs	1.48 1.00	1.39 1.00
50	415709 415735	AA049850 AA704162	Hs.120811	ESTs, Weakly similar to I38022 hypotheti	1.00	72.00
	415799	AA653718	Hs.225841	DKFZP434D193 protein	6.23	31.00
	415817 415857	U88967 AA866115	Hs.78867 Hs.127797	protein tyrosine phosphatase, receptor-t Horno saplens cDNA FLJ11381 fis, clone HE	24.30 32.51	1.00 35.00
85	415989	Al267700		ESTs	78.89	1.00

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	416018	AW138239	Hs.78977	proprotein convertase subtilisin/kexin t	1.00	1.00
	416065	BE267931	Hs.78996	proliferating cell nuclear antigen	3.35	2.32
	416111	AA033813	Hs.79018	chromatin assembly factor 1, subunit A (39.03	3.00
_	416177	AA174069	Hs.187607	ESTs	1.00	9.00
5	416178	AI808527	Hs.192822	serologically defined breast cancer anti ESTs, Weakly similar to MUC2_HUMAN MUCIN	3.83 3.67	3.76 1.00
	416208 416209	AW291168 AA236776	Hs.41295 Hs.79078	MAD2 (mitotic arrest deficient, yeast, h	9.70	1.00
	416239	AL038450	Hs.48948	ESTs	83.87	129.00
	416250	AA581386	Hs.73452	hypothetical protein MGC10791	1.96	2.12
10	416322	BE019494	Hs.79217	pyrroline-5-carboxylate reductase 1	2.08	1.73
	416423	H54375	Hs.268921	ESTs	1.00	89.00
	416448	L13210	Hs.79339	lectin, galactoside-binding, soluble, 3 potassium channel, subfamily K, member 1	1.28 27.29	1.54 67.00
	416498 416658	U33632 U03272	Hs.79351 Hs.79432	fibrillin 2 (congenital contractural ara	53.29	51.00
15	416661	AA634543	Hs.79440	IGF-II mRNA-binding protein 3	9.96	5.00
	416722	AA354604	Hs.122546		3.68	33.00
	416819	U77735	Hs.80205	pim-2 oncogene	1.59	1.84
	416936	N21352	Hs.42987	ESTs, Weakly similar to S21348 probable	1.00	1.00
20	417034	NM_006183	Hs.80962	neurolensin	1.00	1.00 156.00
20	417061 417079	AI675944	Hs.188691 Hs.81134	Homo sapiens cDNA FLJ12033 fis, clone HE Interleukin 1 receptor antagonist	32.95 3.91	4.93
	417079	U65590 AA129547	Hs.285754	met proto-oncogene (hepatocyte growth fa	1.00	51.00
	417233	W25005	Hs.24395	small inducible cytokine subfamily B (Cy	3.38	2.05
0.0	417308	H60720	Hs.81892	KIAA0101 gene product	82.94	25.36
25	417315	AI080042	Hs.180450	ribosomal protein S24	106.61	121.00
	417324	AW265494	N= 4070	ESTS	1.20 8.97	1.28 3.27
	417366 417389	BE185289 BE260964	Hs.1076 Hs.82045	small proline-rich protein 1B (cornifin) midkine (neurite growth-promoting factor	2,59	1.82
	417428	N87579	Hs.278871	gb:LL2030F Human fetal heart, Lambda ZAP	1.00	52.00
30	417433	BE270266	Hs.82128	5T4 encofetal trophoblast glycoprotein	304.75	173.00
	417466	AI681547	Hs.59457	hypothetical protein FLJ22127	1.24	1.34
	417512	A1979168	Hs.344096	glycoprotein (transmembrane) nmb	2.14	5.50
	417515	L24203	Hs.82237	ataxia-telanglectasia group D-associated	2.66 1.28	1.68 1.35
35	417542 417576	J04129 AA339449	Hs.82269 Hs.82285	progestagen-associated endometrial prote phosphoribosylglycinamide formyltransfer	42.76	51.00
55	417715	AW969587	Hs.86366	ESTs	6.35	2.75
	417720	AA205625	Hs.208067	ESTs	113.31	56.00
	417791	AW965339	Hs.111471	ESTs	39.98	16.00
40	417830	AW504786	Hs.122579	hypothetical protein FLJ10461	2.61	31.00 2.44
40	417866 417900	AW067903 BE250127	Hs.82772 Hs.82906	collagen, type XI, aipha 1 CDC20 (cell division cycle 20, S. cerevi	2.35 1.52	1.11
	417933	X02308	Hs.82962	thymidylate synthetase	4.74	2.55
	417944	AU077196	Hs.82985	collagen, type V, alpha 2	3.61	5.21
4 ==	417975	AA641836	Hs.30085	hypothetical protein FLJ23186	12.49	38.00
45	417991	AA731452	Hs.19000B	ESTs	1.00	26.00
	418004	U37519	Hs.87539	aldehyde dehydrogenase 3 family, member	3.02 187.59	2.12 1.00
	418007 418054	M13509 NM_002318	Hs.83169 Hs.83354	matrix metalloproteinase 1 (interstitial lysyl oxidase-like 2	2.85	2.63
	418057	NM_012151	Hs.83363	coagulation factor VIII-associated (intr	1.54	1.69
50	418113	Al272141	Hs.83484	SRY (sex determining region Y)-box 4	6.82	5.22
	418140	BE613836 .	Hs.83551	microfibrillar-associated protein 2	1.26	1.46
	418203	X54942	Hs.83758	CDC28 protein kinase 2	134.19	144.00
	418207 418216	C14685 AA662240	Hs.34772 Hs.283099	ESTs AF15q14 protein	1.00 64.66	1.00 61.00
55	418236	AW994005	Hs.337534	ESTs	18.53	147.00
-	418249	H89226	Hs.34892	KIAA1323 protein	30.53	106.00
	418281	U09550	Hs.1154	oviductal giycoprotein 1, 120kD (mucin 9	1.00	3.00
	418283	S79895	Hs.83942	cathepsin K (pycnodysostosis)	3.96	5.16
60	418300	AI433074	Hs.86682	Homo sapiens cDNA: FLJ21578 fis, clone C	3.18 11.96	2.91 6.6B
00	418322 418327	AA284166 U70370	Hs.84113 Hs.84136	cyclin-dependent kinase inhibitor 3 (CDK paired-like homeodomain transcription fa	11.96 9.23	2.22
	418345	AJ001696	Hs.241407	serine (or cysteine) proteinase inhibito	1.00	1.00
	418379	AA218940	Hs.137516	fidgetin-like 1	21.68	44.00
65	418397	NM_001269	Hs.84746	chromosome condensation 1	1.00	8.00
65	418403	D86978	Hs.84790	KIAA0225 protein	16.91 1.56	18.98 1.16
	418462 418478	BE001596 U38945	Hs.85266 Hs.1174	integrin, beta 4 cyclin-dependent kinase Inhibitor 2A (me	3.22	2.38
	418506	AA084248	Hs.85339	G protein-coupled receptor 39	2.66	2.22
	418526	BE019020	Hs.85838	solute carrier family 16 (monocarboxylic	2.04	2.21
70	418538	BE244323	Hs.85951	exportin, tRNA (nuclear export receptor	1.33	37.00
	418543	NM_005329	Hs.85962	hyaluronan synthase 3	1.04	1.23
	418574 418592	N28754 X99226	Hs.284153	M-phase phosphoprotein 9 Fanconi anemia, complementation group A	48.60 18.24	85.00 26.00
	418641	BE243136	Hs.86947	a disintegrin and metalloproteinase doma	1.19	1.41
75	418661	NM_001949	Hs.1189	E2F transcription factor 3	29.05	43.00
-	418663	AK001100	Hs.41690	desmocollin 3	112.17	19.00
	418678	NM_001327	Hs.87225	cancer/testis antigen	1.18	1.10
	418686	Z36830	Hs.87268	annexin A8	1.54	1.98
80	418689 418712	Al360883 Z42183	Hs.274448	hypothetical protein FLJ11029 gb:HSC0BF041 normalized infant brain cDN	1.19 1.00	1.04 12.00
50	418712	AA227609	Hs.94834	ESTs	1.00	49.00
	418738	AW388633	Hs.6682	solute carrier family 7, (cationic amino	49.85	1.00
	418819	AA228776	Hs.191721	ESTs	1.00	140.00
95	418830	BE513731	Hs.88959	hypothetical protein MGC4816	20.97	23.00
85	418882	NM_004996	Hs.89433	ATP-binding cassette, sub-family C (CFTR	57.09	35.00

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	418971	AA360392	Hs.87113	ESTs	1.00	12.00
	418973	AA233056	Hs.191518 Hs.89584	ESTs insulinoma-associated 1	4.89 1.00	28.00 10.00
	419078 419079	M93119 AW014836	Hs.18844	ESTs	1.09	1.98
5	419080	AW150835	Hs.18878	hypothetical protein FLJ21620	2.06	1.68
	419088	AI538323	Hs.52620	integrin, beta 8	15.60 1.11	51.00 1.83
	419092 419121	J05581 AA374372	Hs.89603 Hs.89626	mucin 1, transmembrane parathyroid hormone-like hormone	1.00	1.00
4.0	419171	NM_002846	Hs.89655	protein tyrosine phosphatase, receptor t	1.10	1.14
10	419183	U60669	Hs.89663	cytochrome P450, subfamily XXIV (vitamin	1.00 3.18	1.00 2.43
	419216 419288	AU076718 AA256106	Hs.164021 Hs.87507	smail inducible cytokine subfamily B (Cy ESTs	1.00	34.00
	419335	AW960146	Hs.284137	hypothetical protein FLJ12888	1.00	8.00
16	419354	M62839	Hs.1252	apolipoprotein H (beta-2-glycoprotein i)	22.63	54.00
15	419359 419423	AL043202 D26488	Hs.90073 Hs.90315	chromosome segregation 1 (yeast homolog) KIAA0007 protein	2.50 1.00	1.98 7.00
	419443	D62703	110.00010	gb:HUM316G10B Clontech human aorta polyA	1.00	12.00
	419452	U33635	Hs.90572	PTK7 protein tyrosine kinase 7	1.64	1.84
20	419474 419485	AW968619 AA489023	Hs.155849 Hs.99807	ESTs ESTs, Weakly similar to unnamed protein	13.63 4.27	62.00 2.26
20	419488	AA316241	Hs.90691	nucleophosmin/nucleoplasmin 3	3.66	3.63
	419502	AU076704		fibrinogen, A alpha polypeptide	13.05	115.00
	419539 419556	AF070590 U29615	Hs.90869 Hs.91093	Homo sapiens clones 24622 and 24623 mRNA chitinase 1 (chitotriosidase)	74.60 1.47	117.00 4.98
25	419569	AI971651	Hs.91143	jagged 1 (Alagille syndrome)	1.00	4.00
	419594	AA013051	Hs.91417	topolsomerase (DNA) II binding protein	94.30	94.00
•	419703 419721	AI793257 NM_001650	Hs.128151 Hs.288650	ESTs aquaporin 4	15.26 1.00	50.00 191.00
	419729	AA586442	Hs.21411	gb:no53a03.s1 NCI_CGAP_SS1 Homo sapians	1.00	59.00
30	419741	NM_007019	Hs.93002	ubiquitin carrier protein E2-C	2.02	1.08
	419745 419752	AF042001 AA249573	Hs.93005 Hs.152618	slug (chicken homolog), zinc finger prot ESTs, Moderately similar to ZN91_HUMAN Z	1.00 29.87	1.00 77.00
	419839	U24577	Hs.93304	phospholipase A2, group VII (platelet-ac	50.99	214.00
25	419936	A1792788	11- 00000	gb:ol91d05.y5 NCI_CGAP_Kid5 Homo sapiens	1.00	1.00 2.47
35	419937 419983	AB040959 W55956	Hs.93836 Hs.94030	DKFZP434N014 protein Homo sapiens mRNA; cDNA DKFZp586E1624 (f	1.64 15.72	94.00
	420005	AW271106	Hs.133294	ESTs	3.15	1.43
	420047	A1478658	Hs.94631	brefeldin A-inhibited guanine nucleotide	12.45	39.00 117.00
40	420058 420162	AK001423 BE378432	Hs.94694 Hs.95577	Homo sapiens cDNA FLJ10561 fis, clone NT cyclin-dependent kinase 4	1.00 1.43	1.21
	420251	AW374968	Hs.348112	Human DNA sequence from clone RP5-1103G7	2.35	3.23
	420259	AF004884	Hs.96253	catclum channel, voltage-dependent, P/Q	0.77 45.04	1.15 54.00
	420281 420309	A1623693 AW043637	Hs.323494 Hs.21766	ESTs ESTs, Weakly similar to ALU5_HUMAN ALU S	49.22	31.00
45	420332	NM_001756	Hs.1305	serine (or cysteine) proteinase inhibito	0.05	2.82
	420380	AA640891	Hs.102406	ESTs	0.99 1.00	2.74 1.00
	420462 420520	AF050147 AK001978	Hs.97932 Hs.98510	chondromodulin I precursor similar to rab11-binding protein	49.74	133.00
	420552	AK000492	Hs.98808	hypothetical protein	94.65	88.00
50	420560	AW207748	Hs.59115	ESTs distal-less homeo box 5	1.00 1.00	17.00 13.00
	420610 420689	Al683183 H79979	Hs.99348 Hs.88678	ESTs	50.09	95.00
	420721	AA927802	Hs.159471	ZAP3 protein	1.00	31.00
55	420759	T11832 Al659838	Hs.127797 Hs.99923	Homo saplens cDNA FLJ11381 fis, clone HE lectin, galactoside-binding, soluble, 7	1.00 3.04	48.00 1.25
55	420783 420900	AL045633	Hs.44269	ESTs	2.24	7.00
	420931	AF044197	Hs.100431	small inducible cytokine B subfamily (Cy	1.00	8.00
	421002 421027	AF116030 AA761198	Hs.100932 Hs.55254	transcription factor 17 ESTs	1.00 2.87	27.00 38.00
60	421037	AI684808	Hs.197653	ESTs	1.00	46.00
	421041	N36914	Hs.14691	ESTs, Moderately similar to I38022 hypot	1.00	98.00
	421073 421110	NM_004689 AJ250717	Hs.101448 Hs.1355	metastasis associated 1 cathepsin E	1.34 119.47	1.46 427.00
~ -	421133	AA401369	Hs.190721	ESTs	1.10	17.00
65	421150	AI913562	Hs.189902	ESTs	1.45 1.00	1.63 15.00
	421155 421307	H87879 BE539976	Hs.102267 Hs.103305	lysyl oxidase Homo saplens mRNA; cDNA DKFZp434B0425 (f	1.37	1.10
	421316	AA287203	Hs.324728	SMA5	1.00	21.00
70	421379	Y15221	Hs.103982	small Inducible cytokine subfamily B (Cy	1.92 5.89	3.94 14.00
70	421451 421474	AA291377 U76362	Hs.50831 Hs.104637	ESTs solute carrier family 1 (glutamate trans	1.46	1.76
	421506	BE302796	Hs.105097	thymidine kinase 1, soluble	1.56	1.08
	421508	NM_004833 Y11339	Hs.105115 Hs.105352	absent in melanoma 2 GalNAc alpha-2, 6-sialyltransferase I, I	5.11 1.00	5.23 3.00
75	421515 421524	AA312082	Hs.105345	GDNF family receptor alpha 1	2.63	10.58
	421526	AL080121	Hs.105460	DKFZP564O0823 protein	1.46	1.88
	421552 421574	AF026692 AJ000152	Hs.105700 Hs.105924	secreted frizzled-related protein 4 defensin, beta 2	30.21 1.67	50.32 1.74
	421574 421582	AJ000152 Al910275	ng. 100324	trefoll factor 1 (breast cancer, estroge	1.23	1.00
80	421633	AF121860	Hs.106260	sorting nexin 10	1.00	116.00
	421659 421677	NM_014459 H64092	Hs.106511 Hs.38282	protocadherin 17 ESTs	0.05 1.31	6.33 1.42
	421753	BE314828	Hs.107911	ATP-binding cassette, sub-family B (MDR/	1.41	1.20
95	421773	W69233	Hs.112457	ESTs	1.12	1.14 1.29
85	421777	BE562088	Hs.108196	HSPC037 protein	1.97	1.23

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	421800	AA298151	Hs.222969	ESTs	1.03	1.30
	421817	AF146074	Hs.108660	ATP-binding cassette, sub-family C (CFTR	1.88 11.84	1.59 22.80
	421896 421928	N62293 AF013758	Hs.45107 Hs.109643	ESTs polyadenylate binding protein-interactin	45.89	90.00
5	421931	NM_000814	Hs.1440	gamma-aminobutyric acid (GABA) A recepto	1.13	1.49
	42194B	L42583	Hs.334309	keratin 6A	51.83	20.25
	421975	AW961017	Hs.6459	hypothetical protein FLJ11856	1.17 1.00	1.15 52.00
	422026 422094	U80736 AF129535	Hs.110826 Hs.272027	trinucleotide repeat containing 9 F-box only protein 5	67.61	62.00
10	422095	AI868872	Hs.282804	hypothetical protein FLJ22704	4.37	2.34
	422109	S73265	Hs.1473	gastrin-releasing peptide	4.18	95.50
	422128	AW881145	11 4470	gb:QV0-OT0033-010400-182-a07 OT0033 Homo	40.89 1.13	71.00 1.38
	422129 422134	AU076635 AW179019	Hs.1478 Hs.112110	serine (or cysteine) proteinase inhibito mitochondrial ribosomal protein L42	41.59	96.00
15	422158	L10343	Hs.112341	protease Inhibitor 3, skin-derived (SKAL	2.37	1.10
	422168	AA586894	Hs.112408	S100 calcium-binding protein A7 (psortas	3.29	1.68
	422278	AF072873	Hs.114218	frizzled (Drosophila) homolog 6	4.93 1.49	5.73 1.71
	422282 422283	AF019225 AW411307	Hs.114309 Hs.114311	apolipoprotein L CDC45 (cell division cycle 45, S.cerevis	25.99	10.91
20	422310	AA316622	Hs.98370	cytochrome P450, subfamily IIS, polypept	1.54	1.41
	422311	AF073515	Hs.114948	cytokine receptor-like factor 1	1.15	1.78
	422330	D30783	Hs.115263	epiregulin	1.00	112.00
	422364 422406	AF067800 AF025441	Hs.115515 Hs.116206	C-type (calcium dependent, carbohydrate- Opa-Interacting protein 5	9.39 18.33	60.00 53.00
25	422424	Al186431	Hs.296638	prostate differentiation factor	1.71	3.21
	422440	NM_004812	Hs.116724	aldo-keto reductase family 1, member B10	47.53	32.00
	422487.	AJ010901	Hs.198267	mucin 4, tracheobronchial	73.68	35.54
	422511 422515	AU076442 AW500470	Hs.117938 Hs.117950	collagen, type XVII, atpha 1 multifunctional polypeptide similar to S	173.97 4.68	26.00 2.92
30	422656	AI870435	Hs.1569	LIM homeobox protein 2	1.00	1.00
	422737	M26939	Hs.119571	collagen, type III, alpha 1 (Ehlers-Dan)	3.89	4.55
•	422756	AA441787	Hs.119689	glycoprotein hormones, alpha polypeptide	1.05 3.88	1.46 1.53
	422765 422809	AW409701 AK001379	Hs.1578 Hs.121028	baculoviral IAP repeat-containing 5 (sur hypothetical protein FLJ 10549	99.56	53.00
35	422867	L32137	Hs.1584	cartilage oligomeric matrix protein (pse	1.69	3.17
	422938	NM_001809	Hs.1594	centromere protein A (17kD)	70.46	61.00
	422956	BE545072	Hs.122579	ECT2 protein (Epithelial cell transformi	77.74 5.88	3.00 8.55
	422960 422963	AW890487 AA401369	Hs.63984 Hs.190721	cadherin 13, H-cadherin (heart) ESTs	171.41	17.00
40	422976	AU076657	Hs.1600	chaperonin containing TCP1, subunit 5 (e	2.12	1.62
	422981	AF026445	Hs.122752	TATA box binding protein (TBP)-associate	10.49	35.00
	422986	AA319777	Hs.221974	ESTS	12.40 16.41	32.47 60.00
	423034 423049	AL119930 X59373	Hs.188023	gb:DKFZp761A092_r1 761 (synonym: hamy2) ESTs, Moderately similar to HXDA_HUMAN H	1.00	1.00
45	423081	AF262992	Hs.123159	sperm associated antigen 4	1.82	2.96
	423184	NM_004428	Hs.1624	ephrin-A1	1.14	1.53
	423217 423248	NM_000094 AA380177	Hs.1640 Hs.125845	collagen, type VII, alpha 1 (epidermolys ribulose-5-phosphate-3-epimerase	2.14 7.18	1.69 14.00
	423309	BE006775	Hs.126782	sushi-repeat protein	21.90	64.00
50	423361	AW170055	Hs.47628	ESTs	1.00	1.00
	423453	AW450737	Hs.128791	CGI-09 protein	55.52 0.88	66.00 1.17
	423511 423516	AF036329 AB007933	Hs.129715 Hs.129729	gonadotropin-releasing hormone 2 ligand of neuronal nitric oxide synthase	1.76	5.40
	423551	AA327598	Hs.233785	ESTs	3.54	4.33
55	423554	M90516	Hs.1674	glutamine-fructose-6-phosphate transamin	1.00	50.00
	423575 423624	C18863 Al807408	Hs.163443 Hs.166368	Homo sapiens cDNA FLJ11576 fis, clone HE ESTs	38.88 1.00	70.00 67.00
	423634	AW959908	Hs.1690	heparin-binding growth factor binding pr	76.02	1.00
	423642	AW452650	Hs.157148	hypothetical protein MGC13204	19.14	58.00
60	423662	AA642452	Hs.130881	B-ceil CLL/lymphoma 11A (zinc finger pro matrix metaltoproteinase 12 (macrophage	3.61 240.73	13.57 40.00
	423673 423698	BE003054 AA329796	Hs.1695 Hs.1098	DKFZp434J1813 protein	1.00	59.00
	423725	AJ403108	Hs.132127	hypothetical protein LOC57822	4.20	1.00
65	423761	NM_006184	Hs.132576	paired box gene 9	1.00	1.00
65	423787 423816	AJ295745 AF151064	Hs.236204	nuclear pore complex protein hypothetical protein	7.16 1.00	6.64 44.00
	423826	U20325	Hs.1707	cocaine- and amphetamine-regulated trans	1.00	1.00
	423849	AL157425	Hs.133315	Homo sapiens mRNA; cDNA DKFZp761J1324 (f	1,00	1.00
70	423887	AL080207	Hs.134585	DKFZP434G232 protein	1.00 31.33	1.00 31.00
70	423934 423954	U89995 AW753164	Hs.159234 Hs.288604	forkhead box E1 (thyroid transcription f KIAA1632 protein	5.81	10.87
	423961	D13666	Hs.136348	osteoblast specific factor 2 (fasciclin	3.55	3.30
	424012	AW368377	Hs.137569	tumor protein 63 kDa with strong homolog	233.42	68.00
75	424016	AW163729	Hs.6140 Hs.153692	hypothetical protein MGC15730 Homo sapiens cDNA FLJ14354 fis, clone Y7	0.93 21.30	1.01 52.00
13	424028 424046	AF055084 AF027866	Hs.138202	serine (or cysteine) proteinase inhibito	1.00	1.00
	424086	Al351010	Hs.102267	lysyl oxidase	21.91	70.00
	424098	AF077374	Hs.139322	small proline-rich protein 3	137.82	54.00
80	424120 424165	T80579 AW582904	Hs.290270 Hs.142255	ESTs islet amylold polypeptide	1.00 1.00	1.00 34.00
-	424200	AA337221	1 10, 142200	gb:EST41944 Endometrial turnor Homo sapie	13.06	48.00
	424279	L29306	Hs.171814	tryptophan hydroxylase (tryptophan 5-mon	1.00	1.00
	424308	AW975531	Hs.154443	minichromosome maintenance deficient (S.	164.58 53.72	87.00 302.00
85	424326 424340	NM_014479 AA339036	Hs.145296 Hs.7033	disintegrin protease ESTs	0.88	1.15
					-	

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	424351	BE622117	Hs.145567	hypothetical protein	0.93	1.03
	424364	AW383226	Hs.201189	ESTs, Weakly similar to G01763 atrophin-	7.02	3.24
	424381	AA285249	Hs.146329	protein kinase Chk2	95.55	92.00
-	424411	NM_005209	Hs.146549	crystallin, beta A2	1.63	3.25
5	424420 424441	BE614743	Hs.146688 Hs.147097	prostaglandin E synthase	1.63 1.82	1.33 1.29
	424502	X14850 AF242388	Hs.147057	H2A histone family, member X lengsin	1.00	1.00
	424503	X06256	Hs.149609	Integrin, alpha 5 (fibronectin receptor,	1.02	2.24
4.0	424513	BE385864	Hs.149894	mitochondrial translational initiation f	1.00	17.00
10	424539	L02911	Hs.150402	Activin A receptor, type I (ACVR1) (ALK	32.46	108.00
	424568	AF005418	Hs.150595	cytochrome P450, subfamily XXVIA, polype	3.40	2.58
	424602	AK002055	Hs.151046	hypothetical protein FLJ11193	31.87	25.00 2.37
	424629 424645	M90656 NM_014682	Hs.151393 Hs.151449	glutamate-cysteine ligase, catalytic sub KIAA0535 gene product	3.58 1.00	1.00
15	424687	J05070	Hs.151738	matrix metalloproteinase 9 (gelatinase B	2.12	2.23
10	424717	AW992292	Hs.152213	wingless-type MMTV integration site fami	1.00	1.00
	424834	AK001432	Hs.153408	Homo sapiens cDNA FLJ10570 fis, clone NT	56.19	12.00
	424840	D79987	Hs.153479	extra spindle poles, S. cerevislae, homo	2.65	1.30
20	424867	AI024860	Hs.153591	Not56 (D. melanogaster)-like protein	1.23	1.05
20	424905	NM_002497	Hs.153704	NIMA (never in mitosis gene a)-related k	21.35	1.00
	424979 424999	D87989 AW953120	Hs.154073	UDP-galactose transporter related gb:EST365190 MAGE resequences, MAGB Homo	1.36 1.24	1.35 1.41
	425048	H05468	Hs.164502	ESTs	1.00	11.00
	425057	AA826434	Hs.1619	achaete-scute complex (Drosophila) homol	7.46	87.00
25	425081	X74794	Hs.154443	minichromosome maintenance deficient (S.	2.52	3.82
	425118	AU076611	Hs.154672	methylene tetrahydrofolate dehydrogenase	4.84	4.03
	425159	NM_004341	Hs.154868	carbamoyl-phosphate synthetase 2, aspart	3.62	2.73
	425202	AW962282	Hs.152049	ESTs, Weakly similar to 138022 hypotheti	1.00 100.77	53.00 44.00
30	425234 425236	AW152225 AW067800	Hs.165909 Hs.155223	ESTs, Weakly similar to 138022 hypotheti stanniocalcin 2	3.30	2.90
50	425245	A1751768	Hs.155314	KIAA0095 gene product	1.91	2.32
	425247	NM_005940	Hs.155324	matrix metalloproteinase 11 (stromelysin	1.41	1.49
	425266	J00077	Hs.155421	alpha-fetoprotein	1.00	68.00
25	425274	BE281191	Hs.155462	minichromosome maintenance deficient (mi	1.97	1.63
35	425322	U63630	Hs.155637	protein kinase, DNA-activated, catalytic	141.49 1.00	123.00 84.00
	425349 425371	AA425234 D49441	Hs.79886 Hs.155981	ribose 5-phosphate isomerase A (ribose 5 mesothelin	0.87	1.59
	425397	J04088	Hs.156346	topoisomerase (DNA) II alpha (170kD)	14.90	5.76
	425420	BE536911	Hs.234545	hypothetical protein NUF2R	1.00	1.00
40	425424	NM_004954	Hs.157199	ELKL motif kinase	10.58	9.74
	425483	AF231022	Hs.158159	FAT turnor suppressor (Drosophile) homolo	1.74	1.40
	425566	AW162943	Hs.250618	UL16 binding protein 2	1.49	1.14 233.00
	425580 425650	L11144 NM 001044	Hs.1907 Hs.1925	galanin desmoglein 3 (pemphigus vulgaris antigen	53.29 33.45	1.00
45	425692	NM_001944 D90041	Hs.155956	N-acetyltransferase 1 (arylamine N-acety	1.00	55.00
	425695	NM_005401	Hs.159238	protein tyrosine phosphatase, non-recept	1.00	10.00
	425734	AF056209	Hs.159396	peptidylglycine alpha-amidating moncoxyg	1.00	41.00
	425776	U25128	Hs.159499	parathyroid hormone receptor 2	1.00	48.00
50	425810	Al923627	Hs.31903	ESTs	27.39	98.00
30	425811 425849	AL039104 Al077288	Hs.159557 Hs.296323	karyopherin alpha 2 (RAG cohort 1, Impor serum/glucocorticold regulated kinase	1.99 71.16	1.58 3.42
	425852	AK001504	Hs.159651	death receptor 6, TNF superfamily member	1.35	1.34
	426067	AA401369	Hs.190721	ESTs	1.01	17.00
	426088	AF038007	Hs.166196	ATPase, Class I, type 8B, member 1	26.26	47.00
55	426215	AW067800	Hs.155223	stanniocalcin 2	1.91	2.90
	426227	U67058	Hs.154299	Human proteinase activated receptor-2 mR	22.40	25.00
	426269 426283	H15302 NM_003937	Hs.168950 Hs.169139	Homo saplens mRNA; cDNA DKFZp566A1046 (f kynureninase (L-kynurenine hydrolase)	1.00 91.39	1.00 229.00
	426329	AL389951	Hs.271623	nucleoporin 50kD	4.34	4.08
60	426427	M86699	Hs.169840	TTK protein kinase	7.02	1.00
	426432	AF001601	Hs.169857	paraoxonase 2	1.16	1.68
	426440	BE382756	Hs.169902	solute carrier femily 2 (facilitated glu	2.59	1.71
	426459	AF151812	Hs.169992	hypothetical 43.2 Kd protein	1.56 20.60	1.66 26.00
65	426471 426496	M22440 D31765	Hs.170009 Hs.170114	transforming growth factor, alpha KIAA0061 protein	9.81	22.00
05	426501	AA401369	Hs.190721	ESTs	19.23	17.00
	426514	BE616633	Hs.170195	bone morphogenetic protein 7 (osteogenic	103.74	41.00
	426536	A1949749	Hs.44441	ESTs	4.65	23.00
70	426572	AB037783	Hs.170623	hypothetical protein FLJ11183	1.00	43.00
70	426682	AV660038	Hs.2056	UDP glycosyltransferase 1 family, polype	160.08	8.00
	426591 426746	NM_006201 J03626	Hs.171834 Hs.2057	PCTAIRE protein kinase 1 uridine monophosphate synthetase (crotat	1.51 2.13	1.35 1.68
	426752	X69490	Hs.172004	titin	0.02	5.14
	426784	U03749	Hs.172216	chromogranin A (parathyroid secretory pr	1.72	1.71
75	426807	AA385315	Hs.156682	ESTs	1.30	1.64
	426812	AF105365	Hs.172613	solute carrier family 12 (potassium/chlo	1.47	1.53
	426814	AF036943	Hs.172619	myelin transcription factor 1-like	1.00	1.00
	426831 426897	BE296216	Hs.172673	S-adenosylhomocysteine hydrolase ESTs	1.51 141.56	1.25 17.00
80	426925	AA401369 NM_001196	Hs.190721 Hs.315689	Homo sapiens cDNA: FLJ22373 fis, clone H	32.61	38.00
-5	426935	NM_000088	Hs.172928	collagen, type I, alpha 1	2.65	3.16
	426964	AA393739	Hs.287416	Homo saplens cDNA FLJ11439 fis, clone HE	1.97	3.49
	426966	Al493134		sclerostin	1.00	1.00
85	426991	AK001536	Un 470000	Homo sapiens cDNA FLJ10674 fis, clone NT	3.39 4.24	2.28 17.00
95	427099	AB032953	Hs.173560	odd Oz/ten-m homolog 2 (Drosophila, mous	7.47	11.00

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	427239	BE270447	Hs.174070	ubiquitin carrier protein	1.58	1.05
	427260	AA663848	1,0,,,,	gb:ae70b06.s1 Stratagene schizo brain S1	1.34	1.60
	427281	AA906147	Hs.102869	ESTs	1.00	66.00
-	427335	AA448542	Hs.251677	G antigen 7B	51.83	4.00
5	427354	T57896	Hs.191095	ESTs	1.17	1.95
	427356	AW023482	Hs.97849	ESTS	7.31 1.00	41.00 57.00
	427376 427383	AA401533 NM_005411	Hs.19440 Hs.177582	ESTs surfactant, pulmonary-associated protein	0.42	1.32
	427427	AF077345	Hs.177936	lectin, superfamily member 1 (cartilage-	1.00	20.00
10	427441	AA412605	Hs.343879	SPANX family, member C	1.00	1.00
	427445	X80818	Hs.178078	glutamate receptor, metabotropic 4	0.97	1.03
	427505	AA361562	Hs.178761	26S proteasome-associated pad1 homolog	4.60	4.04
	427510	Z47542	Hs.179312	small nuclear RNA activating complex, po	22.00	45.00
15	427528	AU077143	Hs.179565	minichromosome maintenance deficient (S.	97.45	92.00
15	427546	AA188763	Hs.36793 Hs.26534	hypothetical protein FLJ23188 ESTs	1.50 6.81	3.24 40.00
	427562 427585	R56424 D31152	Hs.179729	collagen, type X, alpha 1 (Schmid metaph	69.91	62.00
	427660	Al741320	Hs.114121	Homo sapiens cDNA: FLJ23228 fis, clone C	2.70	49.00
	427666	A1791495	Hs.180142	calmodulin-like skin protein	1.37	1.88
20	427668	AA298760	Hs.180191	hypothetical protein FLJ14904	29.55	67.00
	427677	NM_007045	Hs.180296	FGFR1 oncogene partner	3.52	2.63
	427701	AA411101	Hs.243886	nuclear autoantigenic sperm protein (his	7.41	34.00
	427711	M31659	Hs.180408	solute carrier family 25 (mitochondrial	15.84 7.03	70.00 4.52
25	427719 427722	Al393122 AK000123	Hs.134726 Hs.180479	ESTs hypothetical protein FLJ20116	2.92	1.74
23	427747	AW411425	Hs.180655	serine/threonine kinase 12	1.76	1.26
	427912	AL022310	Hs.181097	tumor necrosis factor (ligand) superfami	9.63	59.00
	427961	AW293165	Hs.143134	ESTs	41.97	118.00
20	428004	AA449563	Hs.151393	glutamate-cysteine ligase, catalytic sub	23.82	1.00
30	428023	AL038843	007504	Homo sapiens cDNA: FLJ23602 fis, clone L	1.40	1.33
	428046	AW812795	Hs.337534	ESTs, Moderately similar to 138022 hypot ESTs	96.28 1.25	167.00 1.29
	428093 428098	AW594506 AU077258	Hs.104830 Hs.182429	protein disulfide isomerase-related prot	1.86	1.60
	428129	AI244311	Hs.26912	ESTs	1.00	42.00
35	428169	Al928984	Hs.182793	golgi phosphoprotein 2	2.76	2.11
	428182	BE386042	Hs.293317	ESTs, Weakly similar to GGC1_HUMAN G ANT	1.00	1.00
	428227	AA321649	Hs.2248	small inducible cytokine subfamily B (Cy	85.59	181.00
	428242	H55709	Hs.2250	leukemia inhibitory factor (cholinergic	8.57	21.64
40	428330	L22524	Hs.2256	matrix metalloproteinase 7 (matrilysin,	7.77 0.58	15.90 1.43
40	428434	A1909935	Hs.65551 Hs.184339	Homo sapiens, Similar to DNA segment, Ch	237.53	204.00
	428450 428471	NM_014791 X57348	Hs.184510	KIAA0175 gene product stratifin	6.00	4.60
	428479	Y00272	Hs.334562	cell division cycle 2, G1 to S and G2 to	58.54	16.00
	428484	AF104032	Hs.184601	solute carrier family 7 (cationic amino	3.53	2.15
45	428505	AL035461	Hs.2281	chromogranin B (secretogranin 1)	1.00	1.00
	428532	AF157326	Hs.184786	TBP-interacting protein	1.00	58.00
	428645	AA431400	Hs.98729	ESTs, Weakly similar to 2017205A dihydro	1.00 1.00	16.00 1.00
	428664 428698	AK001666 AA852773	Hs.189095 Hs.334838	similar to SALL1 (sal (Drosophila)-like KIAA1866 protein	187.37	255.00
50	428728	NM_016625	Hs.191381	hypothetical protein	47.24	80.00
-	428748	AW593206	Hs.98785	Ksp37 protein	1.00	87.00
	428758	AA433988	Hs.98502	hypothetical protein FLJ14303	1.06	1.13
	428771	AB028992	Hs.193143	KIAA1069 protein	1.98	92.00
<i>E E</i>	428801	AW277121	Hs.254881	ESTs	1.67	6.15
55	428810	AF068236	Hs.193788	nitric oxide synthase 2A (inducible, hep	1.03 124.17	1.27 43.00
	428839 428845	A1767756 AL157579	Hs.82302 Hs.153610	Homo sapiens cDNA FLJ14814 fis, clone NT KIAA0751 gene product	1.00	1.00
	428959	AF100779	Hs.194680	WNT1 inducible signaling pathway protein	15.16	27.00
	428969	AF120274	Hs.194689	artemin	1.36	1.24
60	429038	AL023513	Hs.194766	seizure related gene 6 (mouse)-like	0.97	3.31
	429065	Al753247	Hs.29643	Homo sapiens cDNA FLJ13103 fis, clone NT	6.82	16.47
	429164	A1688663	Hs.116586	EST8	19.08	67.00
	429170	NM_001394	Hs.2359	dual specificity phosphatase 4	16.18 79.72	105.00 104.00
65	429183 429201	AB014604 X03178	Hs.197955 Hs.198246	KIAA0704 protein group-specific component (vitamin D bind	1.00	1.00
03	429211	AF052693	Hs.198249	gap junction protein, beta 5 (connexin 3	1.33	1.09
	429220	AW207206	110.700240	ESTs	1.00	7.00
	429228	AI553633	Hs.326447	ESTs	39.47	29.25
~ ^	429259	AA420450	Hs.292911	ESTs, Highly similar to S60712 band-6-pr	2.01	1.18
70	429263	AA019004	Hs.198396	ATP-binding cassette, sub-family A (ABC1	1.07	1.00
	429276	AF056085	Hs.198612	G protein-coupled receptor 51	3.70	142.00
	429359	W00482	Hs.2399	matrix metalloproteinase 14 (membrane-in POU domain, class 2, associating factor	1.30 94.09	1.94 86.00
	429412 429413	NM_006235 NM_014058	Hs.2407 Hs.201877	DESC1 protein	41.91	10.00
75	429486	AF155827	Hs.203963	hypothetical protein FLJ10339	12.19	1.00
	429504	X99133	Hs.204238	lipocalin 2 (oncogene 24p3)	1.61	1.08
	429538	BE182592	Hs.11261	small proline-rich protein 2A	4.43	2.90
	429547	AA401369	Hs.190721	ESTs	1.06	17.00
90	429551	AW450624	Hs.220931	ESTs	2.89	65.00
80	429563	BE619413	Hs.2437	eukaryotic translation initiation factor a disintegrin and metalloproteinase doma	1.49 61.86	1.37 100.00
	429597 429610	NM_003816 AB024937	Hs.2442 Hs.211092	LUNX protein; PLUNC (palate lung and nas	1.59	1.69
	429612	AF062649	Hs.252587	pituitary tumor-transforming 1	2.78	1.74
	429616	Al982722	Hs.120845	ESTs	1.00	1.00
85	429656	X05608	Hs.211584	neurofilament, light polypeptide (68kD)	1.00	4.00

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	429663	M68874	Hs.211587	phospholipase A2, group IVA (cytosolic,	69.95	104.00
	429736	AF125304	Hs.212680	tumor necrosis factor receptor superfami	1.25	1.21
	429782	NM_005754	Hs.220689	Ras-GTPase-activating protein SH3-domain	1.00	7.00
5	429903	AL134197	Hs.93597	cyclin-dependent kinase 5, regulatory su	11.80 1.00	1.00 78.00
5	429918 429978	AW873986 AA249027	Hs.119383	ESTs ribosomal protein S6	1.98	3.09
	429986	AF092047	Hs.227277	sine oculis homeobox (Drosophila) homolo	1.00	48.00
	430044	AA464510	Hs.152812	ESTs	69.27	59.00
	430114	AA847744	Hs.99640	ESTs	1.00	1.00
10	430134	BE380149	Hs.105223	ESTs, Weakly similar to T33188 hypotheti	1.00	51.00
	430147	R60704	Hs.234434	hairy/enhancer-of-split related with YRP	1.10	2.22
	430287	AW182459	Hs.125759	ESTs, Weakly similar to LEU5_HUMAN LEUKE quanine nucleotide binding protein 4	1.00 3.80	127.00 1.47
	430294 430300	AI538226 U60805	Hs.32976 Hs.238648	oncostatin M receptor	1.00	35.00
15	430305	NM_004293	Hs.239147	guanine deaminase	92.31	28.00
15	430337	M36707	Hs.239600	calmodulin-like 3	1.18	1.08
	430378	Z29572	Hs.2556	tumor necrosis factor receptor superfami	5.28	66.00
	430388	AA356923	Hs.240770	nuclear cap binding protein subunit 2, 2	16.76	38.00
20	430393	BE185030	Hs.241305	estrogen-responsive B box protein	1.63 1.00	1.50 1.00
20	430439	AL133561	Un 207020	DKFZP434B061 protein	1.64	2.12
	430451 430454	AA836472 AW469011	Hs.297939 Hs.105635	cathepsin B ESTs	63.35	44.00
	430466	AF052573	Hs.241517	polymerase (DNA directed), theta	2.47	1.91
	430481	AA479678	Hs.203269	ESTs, Moderately similar to ALU8_HUMAN A	1.00	31.00
25	430486	BE062109	Hs.241551	chloride channel, calcium activated, fam	12.28	41.00
	430508	Al015435	Hs.104637	ESTs	4.75	7.27
	430533	AA480895	Hs.57749	ESTs, Weakly similar to T17288 hypotheti	1.00 1.00	1.00 1.59
	430563	AF146074	Hs.108660	ATP-binding cassette, sub-family C (CFTR	1.72	1.39
30	430677 430678	Z26317 AA401369	Hs.94560 Hs.190721	desmoglein 2 ESTs	0.90	17.00
50	430686	NM_001942	Hs.2633	desmoglein 1	1.00	1.00
	430788	Al742925	Hs.7179	ESTs, Weakly similar to 2004399A chromos	1.62	1.84
	430890	X54232	Hs.2699	glypican 1	1.58	1.40
25	430935	AW072916		zinc finger protein 131 (clone pHZ-10)	90.28	132.00
35	430985	AA490232	Hs.27323	ESTs, Weakly similar to 178885 serine/th	0.94 60.25	1.28 28.00
	431009 431089	BE149762 BE041395	Hs.48956	gap junction protein, beta 6 (connextn 3 ESTs, Wealdy similar to unknown protein	23.32	941.00
	431092	Al332764	Hs.125757	ESTS	13.46	63.00
	431124	AF284221	Hs.59506	doublesex and mab-3 related transcriptio	49.43	62.00
40	431164	AA493650	Hs.94367	Homo saplens cDNA: FLJ23494 fls, clone L	0.44	2.20
	431211	M86849	Hs.323733	gap junction protein, beta 2, 26kD (conn	182.26	101.00
	431221	AW207837	Hs.286145	SRB7 (suppressor of RNA polymerase B, ye	4.15 1.00	13.97 86.00
	431277 431322	AA501806	Hs.345824	ESTs gb:EST382704 MAGE resequences, MAGK Homo	40.55	200.00
45	431342	AW970622 AW971018	Hs.21659	ESTs	1.00	53.00
	431384	BE158000	Hs.285026	gb:MR2-HT0377-150200-202-e03 HT0377 Homo	0.94	1.14
	431462	AW583672	Hs.256311	granîn-like neuroendocrine peptide precu	1.30	1.25
	431494	AA991355	Hs.298312	hypothetical protein DKFZp434A1315	3.90	26.00
50	431515	NM_012152	Hs.258583	endothelial differentiation, lysophospha	1.41 5.66	1.87 15.00
50	431548 431630	AI834273 NM_002204	Hs.9711 Hs.265829	novel protein integrin, alpha 3 (antigen CD49C, alpha	0.99	1,44
	431745	AW972448	Hs.163425	ESTs	0.99	3.51
	431770	BE221880	Hs.268555	5-3' exoribonuclease 2	67.12	91.00
	431830	Y16645	Hs.271387	smail inducible cytokine subfamily A (Cy	3.36	4.71
55	431846	BE019924	Hs.271580	uroplakin 1B	4.49	2.51
	431890	X17033	Hs.271986	integrin, alpha 2 (CD49B, alpha 2 subuni	2.20 1.01	3.32 1.04
	431934 431958	AB031481 X63629	Hs.272214 Hs.2877	STG protein cadherin (placenta	51.17	46.35
	431936	AL137382	Hs.272320	Homo sapiens mRNA; cDNA DKFZp434L1226 (f	0.94	1.65
60	432023	R43020	Hs.236223	EST	0.94	47.00
	432201	Al538613	Hs.298241	Transmembrane protease, serine 3	1.10	2.24
	432210	Al567421	Hs.273330	Homo sapiens, clone IMAGE:3544662, mRNA,	1.42	1.45
	432226	AW182766	Hs.273558	phosphate cytidylyltransferase 1, cholin	1.00 18.67	1.00 1.00
65	432239	X81334	Hs.2936	matrix metalloproteinase 13 (collagenase SCG10-like-protein	1.09	1.21
05	432265 432281	BE382679 AK001239	Hs.285753 Hs.274263	hypothetical protein FLJ10377	40.98	58.00
	432365	AK001106	Hs.274419	hypothetical protein FLJ10244	1.00	214.00
	432374	W68815	Hs.301885	Homo sapiens cDNA FLJ11346 fis, clone PL	157.34	37.00
~	432375	BE536069	Hs.2962	S100 calcium-binding protein P	1.65	1.06
70	432407	AA221036	11 400404	gb:zr03f12.r1 Stratagene NT2 neuronal pr	73.71	75.00
	432441	AW292425	Hs.163484 Hs.207530	ESTs ESTs	56.35 1.00	72.00 24.00
	432489 432543	A1804855 AA552690	Hs.152423	Homo sapiens cDNA: FLJ21274 fis, clone C	137.72	98.00
	432552	Al537170	Hs.173725	ESTs, Weakly similar to ALU8_HUMAN ALU S	1.00	31.00
75	432583	AW023624	Hs.162282	potassium channel TASK-4; potassium chan	0.27	35.18
-	432606	NM_002104	Hs.3066	granzyme K (serine protease, granzyme 3;	2.87	6.22
	432625	AJ243596	Hs.94830	ESTs, Moderately similar to T03094 A-kin	26.63	56.00
	432653	N62096	Hs.293185	ESTs, Weakly similar to JC7328 amino aci	1.92	5.29 48.00
80	432677	NM_004482	Hs.278611	UDP-N-acetyl-alpha-D-galactosamine:polyp ESTs, Weakly similar to KIAA1074 protein	1.00 45.13	31.00
30	432715 432753	AA247152 NM_014075	Hs.200483 Hs.336938	Homo sapiens PR00593 mRNA, complete cds	1.00	68.00
	432788	AA521091	Hs.178499	Homo saplens cDNA: FLJ23117 ffs, clone L	2.69	3.67
	432842	AW674093	Hs.334822	hypothetical protein MGC4485	1.22	1.34
0.5	432867	AW016936	Hs.233364	ESTs	1.00	1.00
85	432917	NM_014125	Hs.241517	PRO0327 protein	10.25	6.62

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	432920	U37689	Hs.3128	polymerase (RNA) II (DNA directed) polyp	1.44	1.30
	433001	AF217513	Hs.279905 Hs.87409	clone HQ0310 PRO0310p1 thrombospondin 1	154.79 20.96	85.64 100.00
	433023 433042	AW864793 AW193534	Hs.281895	Homo sapiens cDNA FLJ11660 fis, clone HE	1.00	10.00
5	433091	Y12642	Hs.3185	lymphocyte antigen 6 complex, locus D	1.20	1.09
	433159	AB035898	Hs.150587	kinesin-like protein 2	13.82	39.00
	433183	AF231338	Hs.222024	transcription factor BMAL2 ESTs, Weakly similar to ALUB_HUMAN !!!!	1.00 1.00	69.00 1.25
	433258 433409	AA622788 Al278802	Hs.203613 Hs.25661	ESTS	44.81	117.00
10	433437	U20536	Hs.3280	caspase 6, apoptosis-related cysteine pr	70.39	105.00
	433485	Al493076	Hs.201967	aldo-keto reductase family 1, member C2	11.55	2.00
	433537	Al733692	Hs.112488	ESTs	8.66 25.46	55.00
	433547	W04978	Hs.303023	beta tubulin 1, class VI calcium/calmodulin-dependent protein kin	25.16 1.00	83.00 19.00
15	433556 433647	W56321 AA603367	Hs.111460 Hs.222294	ESTs	20.30	49.00
1.5	433658	L03678	Hs.156110	Immunoglobulin kappa constant	5.92	10.03
	433800	AI094221	Hs.135150	lung type-I cell membrane-associated gly	2.29	2.22
	433819	AW511097	Hs.112765	ESTs	3.71 62.08	8.00 104.00
20	433862 433980	D86960 AA137152	Hs.3610 Hs.286049	KIAA0205 gene product phosphoserine aminotransferase	108,91	47.00
20	434088	AF116677	Hs.249270	hypothetical protein PRO1966	1.00	1.00
	434094	AA305599	Hs.238205	hypothetical protein PRO2013	121.27	87.00
	434105	AW952124	Hs.13094	presentlins associated rhombold-like pro	1.22 14.11	1.23 57.00
25	434217 434340	AW014795 Al193043	Hs.23349 Hs.128685	ESTs ESTs, Wealdy similar to T17226 hypotheti	2.10	2.56
23	434360	AA401369	Hs.190721	ESTs	40.98	17.00
	434414	A1798376		gb:tr34b07.x1 NCI_CGAP_Ov23 Homo sapiens	1.48	1.56
	434424	AI811202	Hs.325335	Homo sapiens cDNA: FLJ23523 fis, clone L	1.00	64.00
30	434467 434551	BE552368	Hs.231853 Hs.280858	Homo sapiens cDNA FLJ 13445 ffs, clone PL ESTs, Highly similar to A35661 DNA excls	54.91 2.46	85.00 2.00
50	434627	BE387162 Al221894	Hs.39311	ESTs .	1.00	1.00
	434699	AA643687	Hs.149425	Homo sapiens cDNA FLJ11980 fis, clone HE	1.00	23.00
	434769	AA648884	Hs.134278	Homo sapiens cDNA FLJ12676 fis, clone NT	7.08	56.00
35	434792	AA649253	Hs.132458 Hs.256150	ESTs Homo sapiens, Similar to RIKEN cDNA 2810	8.52 11.33	44.00 1.00
55	434808 434828	AF155108 D90070	Hs.96	phorbol-12-myristate-13-acetate-induced	1.00	1.00
	434876	AF160477	Hs.61460	lg superfamily receptor LNIR	-1.25	1.29
	434891	AA814309	Hs.123583	ESTs	1.00	6.00
40	434928	AW015595	Hs.4267 Hs.110024	Homo saplens clones 24714 and 24715 mRNA Target CAT	1.00 1.26	1.00 1.10
70	435013 435066	H91923 BE261750	Hs.4747	dyskeratosis congenita 1, dyskerin	1.69	1.37
	435087	AW975241	Hs.23567	ESTs	1.00	1.00
	435099	AC004770	Hs.4756	flap structure-specific endonuclease 1	2.90	1.93
45	435159	AA668879	Hs.116649	ESTs	1.00 1.02	1.00 1.46
43	435205 435232	X54136 NM_001262	Hs.181125 Hs.4854	Immunoglobulin lambda locus cyclin-dependent kinase inhibitor 2C (p1	2.04	2.70
	435304	H10709	Hs.269524	ESTs	27.58	139.00
	435313	A1769400	Hs.189729	ESTs	1.00	14.00
50	435505	AF200492	Hs.211238	Interleukin-1 homolog 1	1.00 1.00	38.00 1.00
20	435509 435525	AI458679 AI831297	Hs.181915 Hs.123310	ESTs ESTs	1.00	56.00
	435532	AW291488	Hs.117305	Homo sepiens, clone IMAGE:3682908, mRNA	1.00	2.00
	435550	Al224456	Hs.324507	H.sapiens polyA site DNA	3.42	3.92
55	435602	AF217515	Hs.283532	uncharacterized bone marrow protein BM03	3.95 1.00	1.80 28.00
55	435766 435793	R11673 AB037734	Hs.186498 Hs.4993	ESTs KIAA1313 protein	23.68	42.00
	436069	A1056879	Hs.263209	ESTs	1.00	58.00
	436170	AW450381	Hs.14529	ESTs	1.00	18.00
60	436211	AK001581	Hs.334828	hypothetical protein FLJ10719; KIAA1794	5.84 1.42	22.00 1.27
UU	436213 436217	AA325512 T53925	Hs.71472 Hs.107	hypothetical protein FLJ10774; KIAA1709 fibrinogen-like 1	57.97	31.00
	436238	AK002163	Hs.301724	hypothetical protein FLJ11301	2.51	1.71
	436251	BE515065	Hs.296585	nucleolar protein (KKE/D repeat)	2.33 •	1.64
65	436291	BE568452	Hs.344037	protein regulator of cytokinesis 1 hypothetical protein FLJ23588	108.9 9 0.75	52.00 2.81
UJ	436302 436396	AL355841 AW992292	Hs.99330 Hs.152213	wingless-type MMTV integration site fami	60.01	1.00
	436414	BE264633	Hs.143638	WD repeat domain 4	2.50	2.19
	436419	AI948626	Hs.171356	ESTs	0.95	1.33
70	436443	AW138211	Hs.128746 Hs.199887	ESTs	1.12 1.00	9.26 1.00
70	436474 436481	AJ270693 AA379597	Hs.5199	ESTs HSPC150 protein similar to ubiquitin-con	3.28	1.56
	436486	AA742221	Hs.120633	ESTs	1.00	19.00
	436511	AA721252	Hs.291502	ESTs	16.76	14.00
75	436553	X57809	Hs.181125 Hs.5027	immunoglobulin lambda locus ESTs, Weakly similar to A47582 B-cell gr	1.08 19.20	1.74 9.75
13	436557 436608	W15573 AA628980	ns.ouzi	down syndrome critical region protein DS	33.92	25.00
	436667	AW025183	Hs.127680	ESTs	0.89	1.19
	436771	AW975687	Hs.292979	ESTs	1.00	10.00
80	436839	AA401369 AW953157	Hs.190721 Hs.193235	ESTs hypothetical protein DKFZp547D155	1.00 1.06	17.00 1.15
50	436887 436944	AW268614	Hs.5840	ESTs	1.00	1.00
	436961	AW375974	Hs.156704	ESTs	25.13	25.00
	436972	AA284679	Hs.25640	claudin 3	1.59	1.46
85	437016	AU076916	Hs.5398 Hs.69517	guanine monphosphate synthetase cDNA for differentially expressed CO16 g	2.35 1.34	1.78 1.13
J	437044	AL035864	1100011	and a superior of the second of the A	****	

437161 AJ051515 http://dx.doi.org/10.1001/10.100171.		w	O 02/086	443			
437204					ESTs, Weakly similar to KIAA0758 protein	1.00	17.00
1,00							
State							
43727 AL15746 h. 28846 homo sapiers mRN-C-CDN DR-256560134 (r) 113.25 125.00	5						
437370 AL35567 Hs. 161967 Hs. 162967	3						
437393 Al129559							
43742 5E059288 Hs.34744 Homo sapies mRNA- CDNA DIFEZ-647C135 (fr							
43744 H4608 h153151 ESTs 1.00 130.00 43762 B36800 h153718 h156135 eSTs chromosome condensation-related SMC-asso 1.00 19.00 43762 A3881344 h152781 ESTs, Weakly similar to T17330 hypothet 1.00 3.00 437814 A388192 h153847 ESTs, Weakly similar to T17330 hypothet 1.00 3.00 437916 ESC8028 h15384 h152893 h15384 eSTs, Weakly similar to DX9_HVMAVATP-D 1.00 45.00 437916 ESC8028 h15584 h152893 h152					Homo sapiens mRNA; cDNA DKFZp547C136 (fr		
1979 1979 1970	10						
1.5 1.57 1							
15 43784 A088942							
1.5 337814 A058932 Hz. 135874 EST5, Weakly similar to DDX9_HUMAN ATP-D 1.00 45.00 45785 BEC01839 Hz. 256919 Hz. 25691 Hz. 256919 Hz.							
437840 AA864358 Hs. 256917 45781 457894 Hs. 256987 457916 ESEGOR2 Hs. 256987 457916 ESEGOR2 Hs. 256987 457916 ESEGOR2 Hs. 256987 457916 ESEGOR2 Hs. 202312 457937 AB197222 Hs. 202312 457937 AB197222 Hs. 202312 457937 AB197222 Hs. 202312 457937 AB197222 Hs. 202312 Hs. 202312 457937 AB197222 Hs. 202312 Hs. 202312 457937 AB197222 Hs. 202312 Hs. 202312 Hs. 202312 Hs. 202312 Hs. 202313 AB197302 Hs. 256987 457937 AB197222 Hs. 202312	15						
437816 BE560249 Hs. 5984						1.07	1.78
Agrif Aligner Hard Har				Hs.256897	ESTs, Weakly similar to dJ365O12.1 [H.sa		
Agric Bessel Hand							
437937 All 17222 hbs 12655 ESTs	20						
A39142 A1882058 Ha. 30752b Ha. 30752	20						
438191 AM973052 he ha 8982 ESTs muclear receptor subfamily 1, group 1, m 1.53 10.88 1.89 1.89 1.80 1							
25				110.001020			
438274 A919806 Hs.55080 ESTs 1.00	~ ~			Hs.8882			
438378 A9870529 h.s.282205 ESTs 1.00 1.00 438494 A9808578 h.s.130183 ESTs 2.05 80.00 438456 AW397204 h.s.128511 ESTs 1.00 313.00 438456 AW397204 h.s.128511 ESTs 1.00 313.00 438724 AW812553 h.s.14670 h.s.54618 ESTs 1.00 34.00 438724 AW812553 h.s.14670 h.s.54618 ESTs 1.00 34.00 438724 AW812553 h.s.14670 h.s.54618 h.s.18471 h.s.14617 h.s.146	25				*		
A38403 A386607 Inc. S292206 Inc.							
Age							
A38546							
438552 AJ245820 Hs.5454 Hs.5414 AJ2476 AJ25851 Hs.14670 AJ2476 AJ25851 AJ2476 AJ25851 AJ2477 AJ25851 AJ2477 AJ25851 AJ2477 AJ25851	30						
438724					type I transmembrane receptor (seizure-r	1.43	
ABBS16							
Main							
43885	35						
A38855 A38856 A401269 Hs. 19497 ESTs ESTs 2241 17.00 438916 A401269 Hs. 1959721 ESTs Williams-Beuren syndrome chromosome regi 1.00 1.00 1.00 1.00 439124 439024 856656 Hs. 155656 Hs. 155056 Hs. 156110 Hs. 26018 Hs.	33						
438998 AA401369							
43900			AA401369		ESTs		
439020	40						
A39024 R96896	40			Hs.135056			
A39128				Un 20272			
45 439146 AW13809 Hs.153089 EST6 1,00 67.00 439146 AW13809 Hs.156110 1.65 1.61 1.61 1.62 1.61 1.62 439265 AL133916 4.6527 4.6							
439225							67.00
139285 AL133916 A39318 AW837046 Hs.6527 Hs.114611 A39343 AR085161 Hs.114611 A39451 AF086270 Hs.278554 A39451 AF086270 Hs.278554 A39452 AA918317 Hs.57837 A39453 B254974 Hs.6566 A39472 A99492 AF086310 Hs.103159 A39523 W72348 Hs.103159 ESTs A39502 AF086310 Hs.58551 A39502 A79722	45				immunoglobulin kappa constant		
A39318 AW837046 Hs. 5527 G protein-coupled receptor 56 2.00 2.20				Hs.250618			
A39343 AF086161				U. CE07			
\$\frac{439394}{439410}							
439410 AA632012 Hs.188746 ESTs L83 3.07	50						
12.00					ESTs		
1.58							
55 439477							
A39492	55						
439523 W72348	55						
A39592						1.00	1.19
A39670							
439702 AW085525	C O	439606	W79123	Hs.58561	G protein-coupled receptor 87		
Ay9706 AW872527 Hs.59761 ESTs, Weakly similar to DAP1_HUMAN DEATH 86.55 11.00	00				ESTS, Weakly Similar to ACUU4858 3 U1 SM		
## 439738 BE246502 Hs.9598 sema domain, immunoglobulin domain (lg), 2.36 1.88 439750 AL359053 Hs.57664 Horno sapiens mRNA full length insert CON 2.02 6.08 439750 AL109588 gb:Horno sapiens mRNA full length insert CON 1.00 27.00 439780 AL109588 gb:Horno sapiens mRNA full length insert CON 1.00 27.00 439840 AW449211 Hs.105445 GDNF family receptor alpha 1 1.00 1.00 1.00 439926 AW014875 Hs.137007 ESTs 32.58 71.00 4399379 AW600291 Hs.6823 hypothetical protein FL110430 68.83 61.00 440028 AW473675 Hs.125843 hypothetical protein FL120510 1.83 4.02 440028 AW473675 Hs.125843 ESTs, Weakly similar to T17227 hypothetil 1.42 2.54 440166 AA864968 Hs.127669 KIAA 1603 protein 1.00 54.00 440138 AB033023 Hs.318127 hypothetical protein FL10201 24.18 52.00 440289 AW450991 Hs.192071 ESTs 38.63 113.00 440325 NM_003812 Hs.21433 hypothetical protein FL10201 24.18 52.00 440829 AW450991 Hs.192071 ESTs 38.63 113.00 440829 AV657117 Hs.184164 ESTs, Moderately similar to S65657 alpha 10.84 57.00 440849 AV68298 Hs.146161 hs.7327 daudin 1 3.18 2.37 440704 M69241 Hs.152 insulin-like growth factor binding prote 2.89 2.09 440943 AW082298 Hs.148161 hypothetical protein MGC2408 12.9 17.00 4401001 Hs.272068 ESTs 1.29 17.04 142.99 17.00 142.99 17.0							
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A3980							
1.00 1.00	65	439759		Hs.67709			
100 100				11			
100 100							
70 439979 AW600291 AW600291 Hs.6823 Hs.6824 A40006 AK000517 Hs.6844 hypothetical protein FLJ10430 hppothetical protein FLJ20510 hppothetical prote							
A40026	70						
A40106							
75 440138 AB033023 AB033023 AB033023 Hs.318127 Hypothetical protein FLJ10201 24.18 S2.00 AB05302 AB05302 Hs.32535 Home sapiens cDNA: FLJ23523 fis, clone L S.TS B.G.							
75							
80 AW450991 Hs.192071 Hs.7164 a disintegrin and metalloproteinase doma 62.88 147.00 hypothetical protein DKFZp547J036 2.35 3.62 440527 AV65717 Hs.184164 ESTs, Moderately similar to S65667 alpha 10.84 57.00 440659 AF134160 Hs.7327 claudin 1 3.18 2.37 440704 M69241 Hs.162 Insulin-like growth factor binding prote 2.89 2.09 440943 AW082298 Hs.146161 hypothetical protein MGC2408 2.02 1.41 440994 Al160011 Hs.272068 ESTs 1.29 1.104 441020 AA401369 Hs.190721 ESTs 142.99 17.00	75						
80	, ,						
80 440492 R39127 Hs. 21433 hypothetical protein DKFZp547J036 2.35 3.62 440527 AV657117 Hs. 184164 ESTs, Moderately similar to S65657 alpha 10.84 57.00 440549 AF134160 Hs. 7327 claudin 1 3.18 2.37 440704 M69241 Hs. 162 insulin-like growth factor binding prote 2.89 2.09 440943 AW082298 Hs. 148161 hypothetical protein MGC2408 2.02 1.41 4401920 AA401369 Hs. 190721 ESTs 1.29 17.00							147.00
80		440492	R39127	Hs.21433	hypothetical protein DKFZp547J036		
440704 M69241 Hs.162 Insulin-like growth factor binding prote 2.89 2.09 440943 AW082298 Hs.146161 hypothetical protein MGC2408 2.02 1.41 440994 Al160011 Hs.272068 ESTs 1.29 1.14 441020 AA401369 Hs.190721 ESTs 142.99 17.00	00						57.00
440943 AW082298 Hs.146161 hypothelical protein MGC2408 2.02 1.41 440994 Al160011 Hs.272068 ESTs 1.29 1.14 441020 AA401369 Hs.190721 ESTs 142.99 17.00	δÜ						
440994 A1160011 Hs.272068 ESTs 1.29 1.14 441020 AA401369 Hs.190721 ESTs 142.99 17.00							
441020 AA401369 Hs 190721 ESTs 142.99 17.00							
85 441031 Al110684 · Hs.7645 fibringen, B beta polypeptide 1.41 99.00	0.5				ESTs	142.99	17.00
	85	441031	Al110684	Hs.7645	fibrinogen, B beta polypeptide	1.41	99.00

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	441128	AA570256	11. 00005	ESTs, Weakly similar to T23273 hypotheti	4.13	3.50
	441290 441362	W27501 BE614410	Hs.89605 Hs.23044	cholinergic receptor, nicotinic, alpha p RAD51 (S. cerevisiae) homolog (E coli Re	1.00 130.23	1.00 43.00
_	441377	BE218239	Hs.202656	ESTs .	22.03	1.00
5	441390	AI692560	Hs.131175	ESTs	3.65	7.70
	441497 441525	R51064 AW241867	Hs.23172 Hs.127728	ESTs ESTs	1.00 1.53	1.00 1.42
,	441553	AA281219	Hs.121296	ESTs	1.89	1.57
	441607	NM_005010	Hs.7912	neuronal cell adhesion molecule	1.47	2.11
10	441633 441636	AW958544 AA081846	Hs.112242 Hs.7921	normal mucosa of esophagus specific 1 Homo saplens mRNA; cDNA DKFZp566E183 (fr	216.22 2.31	363.00 2.05
	441737	X79449	Hs.7957	adenosine deaminase, RNA-specific	1.30	1.49
	441790	AA401369	Hs.190721	ESTs	44.15	17.00
15	441801	AW242799	Hs.86366	ESTs ESTs	1,00 1,00	1.00 122.00
13	441919 441937	Al553802 R41782	Hs.128121 Hs.22279	ESTs	0.86	1.37
	441954	A)744935	Hs.8047	Fanconi anemia, complementation group G	1.48	1.39
	442025	AW887434	Hs.11810	CDA11 protein	1.00 9.92	46.00 45.00
20	442029 442072	AW956698 A1740832	Hs.14456 Hs.12311	neural precursor cell expressed, develop Homo sapiens clone 23570 mRNA sequence	25.05	77.00
	442108	AW452649	Hs.166314	ESTs	3,61	3.14
	442117	AW664964	Hs.128899	ESTS	3.00 1.00	5.49 1.00
	442137 442159	AA977235 AW163390	Hs.128830 Hs.278554	ESTs, Weakly similar to Z192_HUMAN ZINC heterochromatin-like protein 1	1.92	1.66
25	442179	AA983842	Hs.333555	chromosome 2 open reading frame 2	27.22	50.00
	442328	AI952430	Hs.150614	ESTs, Weakly similar to ALU4_HUMAN ALU S	5.00	3.42
	442432 442530	BE093589 A1580830	Hs.38178 Hs.176508	hypothetical protein FLJ23468 Homo saplens cDNA FLJ14712 fis, clone NT	181.59 10.59	76.00 144.00
	442547	AA306997	Hs.217484	ESTs, Weakly similar to ALU1_HUMAN ALU S	109.23	98.00
30	442556	AL137761	Hs.8379	Homo sapiens mRNA; cDNA DKFZp586L2424 (f	1.00	53.00
	442619 442710	AA447492 Al015631	Hs.20183 Hs.23210	ESTs, Weakly similar to AF164793 1 prote ESTs	29.02 1.00	50.00 19.00
	442717	R88362	Hs.180591	ESTs, Weakly similar to T23976 hypotheti	1.00	5.00
25	442875	BE623003	Hs.23625	Homo sapiens clone TCCCTA00142 mRNA sequ	22.85	50.00
35	442914 442932	AW188551 AA457211	Hs.99519 Hs.8858	hypothetical protein FLJ14007 bromodomain adjacent to zinc finger doma	25.33 3.18	82.00 4.41
	442942	AW167087	Hs.131562	ESTs	8.45	64.00
	443068	AI188710		ESTs	1.00	27.00
40	443204	AW205878	Hs.29643 Hs.143655	Homo sapiens cDNA FLJ13103 fis, clone NT ESTs	1.00 12,42	24.00 2.00
40	443211 443247	Al128388 BE614387	Hs.333893	c-Myc target JPO1	128.84	96.00
	443324	R44013	Hs.164225	ESTs	0.02	4.59
	443383	A1792453	Hs.166507	ESTS ESTS	1.00 18.52	47.00 61.00
45	443400 443426	R28424 AF098158	Hs.250648 Hs.9329	chromosome 20 open reading frame 1	4.02	1.75
	443572	AA025610	Hs.9605	cleavage and polyadenylation specific fa	2.98	2.57
	443575	A1078022	Hs.269636 Hs.7645	ESTs, Weakly similar to ALU1_HUMAN ALU S fibrinogen, B beta polypeptide	1.00 1.00	29.00 16.00
	443614 443633	AV655386 AL031290	Hs.9654	similar to pregnancy-associated plasma p	1.00	39.00
50	443648	A1085377	Hs.143610	ESTs	39.81	70.00
	443715	A1583187	Hs.9700	cyclin E1 syntaxin 6	48.74 1.29	7.00 1.30
	443723 443802	Al144442 AW504924	Hs.157144 Hs.9805	KIAA1291 protein	1.75	1.61
<i></i>	443859	NM_013409	Hs.9914	follistatin	1.35	1.13
55	443892 443947	AA401369 W24187	Hs.190721	ESTs gb:zb47f09.r1 Soares_fetal_lung_NbHL19W	1.00 1.33	17.00 1.64
	443991	NM_002250	Hs.10082	potassium intermediate/small conductance	5.71	6.87
	444006	BE395085	Hs.10086	type I transmembrane protein Fn14	1.47	1.92
60	444009 444017	A1380792 U04840	Hs.135104 Hs.214	ESTs neuro-oncological ventral antigen 1	1.00 1.00	77.00 1.00
O	444127	N63620	Hs.13281	ESTs	1.00	29.00
	444129	AW294292	Hs.256212	ESTs	1.00	1.00 7.80
	444279 444371	U62432 BE540274	Hs.89605 Hs.239	cholinergic receptor, nicotinic, alpha p forkhead box M1	0.60 2.91	1.14
65	444378	R41339	Hs.12569	ESTs	1.00	1.00
	444381	BE387335	Hs.283713	ESTs, Weakly similar to S64054 hypotheti	469.00	556.00
	444461 444471	R53734 AB020684	Hs.25978 Hs.11217	ESTs, Weakly similar to 2109260A B cell KIAA0877 protein	12.88 24.91	105.00 90.00
	444489	AJ151010	Hs.157774	ESTs	1.00	111.00
70	444619	BE538082	Hs.8172	ESTs, Moderately similar to A46010 X-lin	1.00	70.00
	444665 444707	BE613126 A1188613	Hs.47783 Hs.41690	B aggressive lymphoma gene desmocollin 3	30.5 6 1.00	139.00 1.00
	444735	BE019923	Hs.243122	hypothetical protein FLJ13057 similar to	77.02	90.00
75	444781	NM_014400	Hs.11950	GPI-anchored metastasis-associated prote	1.57	1.31
75	444783	AK001468 AK001676	Hs.62180 Hs.12457	anilin (Drosophila Scraps homolog), act hypothetical protein FLJ10814	77.55 1.00	2.00 27.00
	445236 445258	AK001676 Al635931	Hs.147613	ESTs	1.00	73.00
	445413	AA151342	Hs.12677	CGI-147 protein	28.14	50.00
80	445417 445443	AX001058 AX653838	Hs.12680 Hs.322971	Homo saplens cDNA FLJ10196 fis, clone HE ESTs	1.81 1.00	2.62 1.00
50	445462	AV653838 AA378776	Hs.288649	hypothetical protein MGC3077	2.09	1.70
	445517	AF208855	Hs.12830	hypothetical protein	1.87	70.00
	445537 445580	AJ245671 AF167572	Hs.12844 Hs.12912	EGF-like-domain, multiple 6 skb1 (S. pombe) homolog	1.71 1.52	2.72 1.34
85	445654	X91247	Hs.13046	thioredoxin reductase 1	1.51	1.52

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	445669	AI570830	Hs.174870	ESTs	10.95 1.00	11.45 1.00
	445818 445873	BE045321 AA250970	Hs.136017 Hs.251946	ESTs poly(A)-binding protein, cytoplasmic 1-i	49.42	54.00
_	445885	Al734009	Hs.127699	KIAA1603 protein	1.00	132.00
5	445898	AF070823	Hs.13423	Homo sapiens clone 24468 mRNA sequence	1.00	1.00
	445903 445932	Al347487 BE046441	Hs.132781 Hs.333555	class I cytokine receptor Homo sapiens clone 24859 mRNA sequence	1.00 2.41	36.00 2.88
	445982	BE410233	Hs.13501	pescadillo (zebrafish) homolog 1, contai	1.60	1.35
10	446078	Al339982	Hs.156061	ESTs	1.00	42.00
10	446102 446157	AW168067 BE270828	Hs.317694 Hs.131740	ESTs Homo sapiens cDNA: FLJ22562 fis, clone H	1.00 1.70	1.00 1.53
	446269	AW263155	Hs.14559	hypothetical protein FLJ10540	73.01	48.00
	446292	AF081497	Hs.279682	Rh type C glycoprotein	1.55	1.26
15	446293 446423	AI420213 AW139655	Hs.149722 Hs.150120	ESTS ESTs	1.00 1.10	2.00 4.19
15	446428	AW082270	Hs.12496	ESTs, Weakly similar to ALU4_HUMAN ALU S	0.53	3.26
	446432	Al377320	Hs.150058	ESTs (488) ES	1,00	5.00
	446528 448574	AU076640 Al310135	Hs.15243 Hs.335933	nucleolar protein 1 (120kD) ESTs	1.36 3.89	1,31 72.00
20	446619	AU076643	Hs.313	secreted phosphoprotein 1 (osteopontin,	32.03	20.23
	446636	AC002563	Hs.15767	citron (rho-Interacting, serine/threonin	4.19	5.07
	446783 446839	AW138343 BE091926	Hs.141867 Hs.16244	ESTs mitotic spindle coiled-coil related prot	2.82 110.28	9.47 28.00
	446849	AU076617	Hs.16251	cleavage and polyadenylation specific fa	3.26	2.94
25	446856	AI814373	Hs.164175	ESTs	6.38	11.30
	446872 446880	X97058 AIB11807	Hs.16362 Hs.108646	pyrimidinergic receptor P2Y, G-protein c Homo saplens cDNA FLJ14934 fis, clone PL	1.98 94.90	2.03 113.00
	446921	AB012113	Hs.16530	small inducible cytokine subfamily A (Cy	1.67	3.90
20	446989	AK001898	Hs.16740	hypothetical protein FLJ11036	2.82 1.00	3,12 170.00
30	447022 447033	AW291223 AJ357412	Hs.157573 Hs.157601	ESTs ESTs	7.15	107.00
	447078	AW885727	Hs.9914	ESTs	47.24	24.00
	447081	Y13896	Hs.17287	potassium inwardly-rectifying channel, s	0.12 0.97	17,88 1,48
35	447131 447149	NM_004585 BE299857	Hs.17466 Hs.326	retinolc acid receptor responder (tazaro TAR (HIV) RNA-binding protein 2	1.24	1.26
-	447153	AA805202	Hs.315562	ESTS	1.00	54.00
	447164	AF026941	Hs.17518	Homo sapiens cig5 mRNA, partial sequence	1.00 3.42	67.00 50.00
	447178 447250	AW594641 A1878909	Hs.192417 Hs.17883	ESTs protein phosphatase 1G (formerly 2C), ma	1.60	1.52
40	447289	AW247017	Hs.36978	melanoma antigen, family A, 3	1.00	1.00
	447342	A1199268	Hs.19322	Homo saplens, Similar to RIKEN cDNA 2010 ESTs, Highly similar to S02392 alpha-2-m	28.63 146,62	1.00 51.00
	447343 447350	AA256641 AI375572	Hs.236894 Hs.172634	ESTs	1.00	12.00
4 =	447377	N27687	Hs.334334	transcription factor AP-2 alpha (activat	2.55	63.00
45	447415 447425	AW937335 Al963747	Hs.28149 Hs.18573	ESTs, Weakly similar to KF3B_HUMAN KINES acylphosphatase 1, erythrocyte (common)	0.91 1.00	1.13 35.00
	447519	U46258	Hs.339665	ESTs	59.89	49.00
	447532	AK000614	Hs.18791	hypothetical protein FLI20507	1.23	1.63
50	447534 447636	AA401369 Y10043	Hs.190721	ESTs high-mobility group (nonhistone chromoso	1.00 1.41	17.00 1.11
-	447688	N87079	Hs.19236	Target CAT	1.00	39.00
	447733	AF157482	Hs.19400	MAD2 (mitotic arrest deficient, yeast, h	1.17 6.47	1.12 5.95
	447769 447802	AW873704 AW593432	Hs.320831 Hs.161455	Homo saplens cDNA FLJ14597 fis, clone NT ESTs	0.73	2.34
55	447850	AB018298	Hs.19822	SEC24 (S. cerevisiae) related gene famil	86.45	116.00
	447924	A1817226	Hs.313413 Hs.20141	ESTs, Weakly similar to T23110 hypotheti similar to S. cerevisiae SSM4	1.00 3.50	1.00 4.27
	447973 448030	AB011169 N30714	Hs.325960	membrane-spanning 4-domains, subfamily A	4.13	142.00
60	448105	Al538613	Hs.298241	Transmembrane protease, serine 3	1.15	2.24
60	448243 448278	AW369771 W07369	Hs.52620 Hs.11782	integrin, beta 8 ESTs	15.84 0.97	1.00 1.90
	448290	AK002107	Hs.20843	Homo saplens cDNA FLJ11245 fis, clone PL	1.00	1.00
	448296	BE622756	Hs.10949	Homo saplens cDNA FLJ14162 fis, clone NT	2.42 1.44	2.17 1.08
65	448357 448390	BE274396 AL035414	Hs.108923 Hs.21068	RAB38, member RAS oncogene family hypothetical protein	1.00	43.00
	448469	AW504732	Hs.21275	hypothetical protein FLJ11011	2.63	2.49
	448569	BE382657	Hs.21486 Hs.106823	signal transducer and activator of trans hypothetical protein MGC14797	1.84 3.29	2.53 46.00
	448663 448672	BE614599 Al955511	Hs.225106	ESTs	1.00	21.00
70	448733	NM_005629	Hs.187958	solute carrier family 6 (neurotransmitte	1.82	1.08
	448741 448757	BE614567 Al366784	Hs.19574 Hs.48820	hypothetical protein MGC5469 TATA box binding protein (TBP)-associate	2.48 23.53	1.92 20.00
	448775	AB025237	Hs.388	nudix (nucleoside diphosphate linked moi	2.34	1.97
75	448826	A1580252	Hs.293246	ESTs, Weakly similar to putative p150 [H	74.07	62.67
75	448830 448844	AL031658 Al581519	Hs.22181 Hs.177164	hypothetical protein dJ310013.3 ESTs	1.37 1.00	1.31 31.00
	448988	Y09763	Hs.22785	gamma-aminobutyric acid (GABA) A recepto	1.84	1.95
	448993	Al471630		KIAA0144 gene product alcohol dehydrogenase 7 (class IV), mu o	1.63 1.00	1.49 1.00
80	449003 449029	X76342 N28989	Hs.389 Hs.22891	solute carrier family 7 (calionic amino	1.97	2.26
	449040	AF040704	Hs.149443	putative tumor suppressor	0.97	1.56
	449048	Z45051	Hs.22920	similar to S68401 (cattle) glucose induc ESTs	27.13 8.33	90.00 44.00
	449053 449054	A1625777 AF148848	Hs.344766 Hs.22934	myoneurin	73.85	104.00
85	449101	AA205847	Hs.23016	G protein-coupled receptor	2.58	27.00

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	449167	T05095	Hs.19597	KIAA1694 protein	1.61	2.36
	449207	AL044222	Hs.23255	nucleoporin 155kD	2.36	1.56
	449228	AJ403107	Hs.148590	protein related with psoriasis	1.15	1.15
5	449230	BE613348	Hs.211579	melanoma cell adhesion molecule	206.65 17.28	151.00 45.00
3	449305 449318	A1638293 AW236021	Hs.78531	gb:tt09b07.x1 NCI_CGAP_GC6 Homo sapiens Homo sapiens, Similar to RIKEN cDNA 5730	26.39	35.00
	449448	D60730	Hs.57471	ESTs	1.00	1.00
	449467	AW205006	Hs.197042	ESTs	1.00	1.00
10	449523	NM_000579	Hs.54443	chemokine (C-C motif) receptor 5	56.80	216.86
10	449722 449976	BE280074 H06350	Hs.23960 Hs.135056	cyclin B1 Human DNA sequence from clone RP5-850E9	150.03 2.16	1.00 2.85
	450001	NM_001044	Hs.406	solute carrier family 6 (neurotransmitte	1.17	1.45
	450098	W27249	Hs.8109	hypothetical protein FLJ21080	1.79	2.38
1.5	450101	AV649989	Hs.24385	Human hbc647 mRNA sequence	1.00	69.00
15	450149	AW969781	Hs.132863	Zic family member 2 (odd-paired Drosophi	1.00 29.85	1.00 34.00
	450193 450221	AI916071 AA328102	Hs.15607 Hs.24641	Homo sapiens Fanconi anemia complementat cytoskeleton associated protein 2	1.00	1.00
	450372	BE218107	Hs.202436	ESTs	1.00	1.00
~~	450375	AA009647	Hs.8850	a disintegrin and metalloproteinase doma	51.26	93.00
20	450447	AF212223	Hs.25010	hypothetical protein P15-2	123.20 1.00	181.00
	450568 450589	AL050078 Al701505	Hs.25159 Hs.202526	Homo saplens cDNA FLJ10784 fis, clone NT ESTs	1,00	19.00 23.00
	450684	AA872605	Hs.25333	interleukin 1 receptor, type II	1.00	100.00
~~	450701	H39960	Hs.288467	Homo sapiens cDNA FLJ12280 fis, clone MA	1.89	1.55
25	450705	U90304	Hs.25351	iroquois homeobox protein 2A (IRX-2A) (1.00	45.00
	450832	AA401369 R49131	Hs.190721 Hs.26267	ESTs ATP-dependant interferon response protei	25.17 90.92	17.00 90,00
	450937 450983	AA305384	Hs.25740	ERO1 (S. cerevisiae)-like	3.33	1.70
	451105	Al761324		gb:wi60b11.x1 NCI_CGAP_Co16 Homo saplens	15.02	124.00
30	451110	AI955040	Hs.265398	ESTs, Weakly similar to transformation-r	1.00	143.00
	451253	H48299	Hs.26126	claudin 10	3.02 1.00	2.2 9 1.00
	451291 451320	R39288 AW498974	Hs.6702	ESTs diacytglycerol kinase, zeta (104kD)	2.92	18.00
	451380	H09280	Hs.13234	ESTS	6.90	6.67
35	451386	AB029006	Hs.26334	spastic paraplegia 4 (autosomal dominant	35.75	72.00
	451437	H24143	Hs.31945	hypothetical protein FLJ11071	1.00 1.83	69.00 2.10
	451462 451524	AK000367 AK001466	Hs.26434 Hs.26516	hypothetical protein FLJ20360 hypothetical protein FLJ10604	1.13	1.07
	451541	BE279383	Hs.26557	plakophilin 3	1.88	1.33
40	451592	AI805416	Hs.213897	ESTS	1.00	1.00
	451635	AA018899	Hs.127179	cryptic gene	1.52 4.95	1.92 17.00
	451743 451806	AA401369 NM_003729	Hs.190721 Hs.27076	ESTs RNA 3'-terminal phosphate cyclase	13.55	31.00
	451807	W52854	113.27070	hypothetical protein FLJ23293 similar to	1.55	35.00
45	451871	AI821005	Hs.118599	ESTs	1.81	2.53
	451952	AL120173	Hs.301663	ESTs	1.00	22.00 2.26
	452012 452046	AA307703 AB018345	Hs.279766 Hs.27657	kinesin family member 4A KIAA0802 protein	3.43 56.59	19.00
	452194	AI894413	Hs.332649	olfactory receptor, family 2, subfamily	1.67	4.09
50	452206	AW340281	Hs.33074	Homo sapiens, clone IMAGE:3606519, mRNA,	9.31	53.00
	452240	AA401369	Hs.190721	ESTS	13.42 39.03	17.00 94.00
	452256 452281	AK000933 T93500	Hs.28661 Hs.28792	Homo sapiens cDNA FLJ10071 fis, clone HE Homo sapiens cDNA FLJ11041 fis, clone PL	153.01	340.00
	452291	AF015592	Hs.28853	CDC7 (cell division cycle 7, S. cerevisi	1.95	23.00
55	452295	BE379936	Hs.28866	programmed cell death 10	42.33	61.00
	452304	AA025386	Hs.61311	ESTs, Weakly similar to \$10590 cysteine	1.17 1.00	2.14 13.00
	452340 452349	NM_002202 AB028944	Hs.505 Hs.29189	ISL1 transcription factor, LIM/homeodoma ATPase, Class VI, type 11A	1.09	1.42
	452367	U71207	Hs.29279	eyes absent (Drosophila) homolog 2	54.49	53.00
60	452401	NM_007115	Hs.29352	tumor necrosis factor, alpha-induced pro	1.00	32.00
	452410	AL133619	15-400400	Homo saplens mRNA; cDNA DKFZp434E2321 (f	1.26 24.47	1.99 35.00
	452461 452571	N78223 W31518	Hs.108106 Hs.34665	transcription factor ESTs	54.61	102.00
	452613	AA461599	Hs.23459	ESTs	1.39	1.32
65	452699	AW295390	Hs.213062	ESTs	1.00	26.00
	452705	H49805	Hs.246005	ESTs	1.00	1.00
	452747 452787	AF160477 AW294022	Hs.61460 Hs.222707	lg superfamily receptor LNIR KIAA1718 protein	112.87 1.00	1.29 1.00
	452795	AW392555	Hs.18878	hypothetical protein FLJ21620	1.00	1.00
70	452823	AB012124	Hs.30696	transcription factor-like 5 (basic helix	7.91	75.00
	452833	BE559681	Hs.30736	KIAA0124 protein	3.16	1.92
	452838 452862	U65011 AA401369	Hs.30743 Hs.190721	preferentially expressed antigan in mala ESTs	174.35 98.26	1.00 17.00
	452865	AW173720	Hs.345805	ESTs, Weakly similar to A47582 B-cell gr	1.55	1.00
75	452934	AA581322	Hs.4213	hypothetical protein MGC16207	1.73	1.19
	452946	X95425	Hs.31092	EphA5	1.00	1.00
	452976	R44214	Hs.101189	ESTs	1.58 1.80	1.98 1.60
	453028 453095	AB006532 AW295660	Hs.31442 Hs.252756	RecQ protein-like 4 ESTs	0.77	1.50
80	453102	NM_007197	Hs.31664	frizzled (Drosophila) homolog 10	1.00	1.00
	453103	Al301052	Hs.153444	ESTs	1.00	1.00
	453120	AA292891	Hs.31773	pregnancy-induced growth inhibitor	1.23 1.00	1.20 83.00
	453153 453160	N53893 Al263307	Hs.24360 Hs.239884	ESTs H2B histone family, member L	1.00	30.00
85	453197	AI916269	Hs.109057	ESTs, Weakly similar to ALU5_HUMAN ALU S	1.00	134.00

	w	O 02/0864	143				PCT/I	JS02/12476
	453210	AL133161	Hs.32360	hypothetical protein FLJ10867	1.69	1.93		
	453240	Al969564	Hs.166254	hypothetical protein DKFZp566i133	1.00	1.00		
	453317 453323	NM_002277 AF034102	Hs.41696 Hs.32951	keratin, heir, acidic, 1 solute carrier family 29 (nucleoside tra	1.19 4.90	1.27 4.11		
5	453331	AI240665	Hs.8850	ESTs	199.42	340.00		
	453392	U23752	Hs.32964	SRY (sex determining region Y)-box 11	1.00	16.00		
	453431	AF094754	Hs.32973	glycine receptor, beta	1.00 3.44	1.00 5.17		
	453439 453459	Al572438 BE047032	Hs.32976 Hs.257789	guanine nucleotide binding protein 4 ESTs	2.84	5.58		
10	453563	AW608906.com		Hs.181163	hypothetical	protein MGC5629	4.58	90.00
	453633	AA357001	Hs.34045	hypothetical protein FLJ20764	1.74	1.60		
•	453775 453830	NM_002916 AA534296	Hs.35120 Hs.20953	replication factor C (activator 1) 4 (37 ESTs	19.49 24.92	1.00 25.00		
	453857	AL080235	Hs.35861	DKFZP586E1621 protein	167.59	66.00		
15	453867	A1929383	Hs.33032	hypothetical protein DKFZp434N185	1.00	39.00		
	453883	A1638516	Hs.347524	cofactor required for Sp1 transcriptiona	1.97 63.89	1.58 20.00		
	453884 453900	AA355925 AW003582	Hs.36232 Hs.226414	KIAA0186 gene product ESTs, Weakly similar to ALU8_HUMAN ALU S	20.41	16.00		
	453922	AF053306	Hs.36708	budding uninhibited by benzlmidazoles 1	7.09	22.00		
20	453941	U39817	Hs.36820	Bloom syndrome	29.75	19.00		
	453964 453968	Al961486 AA847843	Hs.12744 Hs.62711	ESTs Homo sapiens, clone IMAGE:3351295, mRNA	1.00 2.06	1.00 1.81		
	453976	BE463830	Hs.163714	ESTs	3.02	131.00		
0.5	454024	AA993527	Hs.293907	hypothetical protein FLJ23403	1.00	131.00		
25	454034	NM_000691	Hs.575	aldehyde dehydrogenase 3 family, member	1.23 30.63	1.02 171.00		
	454042 454059	T19228 NM_003154	Hs.172572 Hs.37048	hypothetical protein FLJ20093 statherin	1.00	1.00		
	454066	X00356	Hs.37058	calcitonin/calcitonin-related polypeptid	1.01	1.45		
20	454098	W27953	Hs.292911	ESTs, Highly similar to S60712 band-6-pr	1.26	1.11		
30	454241 454417	BE144666 Al244459	Hs.110826	gb:CM2-HT0176-041099-017-c02 HT0176 Homo trinucleotide repeat containing 9	6.33 4.30	5.04 7.82		
	454439	AW819152	Hs.154320	DKFZP56601646 protein	1.00	1.00		
	455175	AW993247		gb:RC2-BN0033-180200-014-h09 BN0033 Homo	13.75	103.00		
35	455601	A1368680	Hs.816	SRY (sex determining region Y)-box 2	206.11 1.00	1.00 1.00		
33	456237 456321	AA203682 NM_001327	Hs.87225	gb:zx52e07.r1 Soares_fetal_liver_spleen_ cancer/testis antigen	1.14	1.10		
	458475	NM_000144	Hs.95998	Friedreich ataxia	1.00	48.00		
	456508	AA502764	Hs.123469	ESTs, Weakly similar to AF208855 1 BM-01	162.25	189.00		
40	456534 456736	X91195 AW248217	Hs.100623 Hs.1619	phospholipase C, beta 3, neighbor pseudo achaete-scute complex (Drosophila) homol	2.12 1.15	1.80 1.94		
40	456759	BE259150	Hs.127792	delta (Drosophila)-like 3	1.00	1.00		
	456990	NM_004504	Hs.171545	HIV-1 Rev binding protein	16.42	84.00		
	457200	U33749	Hs.197764 Hs.14355	thyroid transcription factor 1 Homo saplens cDNA FLJ13207 fis, clone NT	0.57 2.71	1.76 · 4.15		,
45	457234 457465	AW968360 AW301344	Hs.122908	DNA replication factor	46.37	47.00		
	457489	A1693815	Hs.127179	cryptic gene	1.12	1.35		
	457646	AA725650	Hs.112948	ESTs	1.55 1.00	2.51 55.00		
	457733 457819	AW974812 AA057484	Hs.291971 Hs.35406	ESTs ESTs, Highly similar to unnamed protein	4.36	3.18		
50	458092	BE545684	Hs.343566	KIAA0251 protein	1.00	1.32		
	458098	BE550224	11- 7000	metallothionein 1E (functional)	1.00	22.00 1.88		
	458207 458242	T28472 BE299588	Hs.7655 Hs.28465	U2 small nuclear ribonucleoprotein auxil Homo saplens cDNA: FLJ21869 fls, clone H	2.06 1.00	1.00		
	458247	R14439	Hs.209194	ESTs	7.00	9.85		
55	458679	AW975460	Hs.142913	ESTs	1.00	3.00		
	458778 458933	AW451034 Al638429	Hs.326525 Hs.24763	arylsulfatase D RAN binding protein 1	1.31 1.98	2.01 1.71		
	459352	AW810383	Hs.206828	ESTs	12.60	63.00		
60	459670	F01020	Hs.172004	tiën	1.00	1.00		
60	459702	A1204995		gb:an03c03.x1 Stratagene schizo brain S1	1.00	237.00		
	TABLE 9	8						
65	Discour	Dalaua Fas	anahanat ida	ettiles sumbre				
05	Pkey: CAT num	ber: Gene clustr		ntifler number	•			
	Accessio		ccession num	bers				
	~ 1	OAT Wb.		t				
70	Pkey 407746	CAT Number 10125_1	Access	son 962 R69415 BE464605 AA418699 AA053293 AA14907	5 AA058396 AW33	8226 AW272659 AA45	4607 AI139535	AW469852 Al275461
, ,	701170	10120_1	AW271	982 AA730033 AA576507 AA991217 AA782067 AI985	851 AA805864 AA	505598 AW469857 R6	9546 AA988279 A	AW001647 N63320
			D8266	1 T27343 AA306950 AA360989 R58778				
	408070	1036688_1		1852 BE35089 5 775 AAD56342 Al538978 AW975281 AA664986				
75	408660 409522	107294_1 113735_1		382 AA075431				
	409866	1156522_1	AW502	2152 H41202 H29772				
	410032	1170435_1		985 BE065944 BE066008 BE066083 BE066093	91 AAGN4004 AACT	007C AAEE440C A400	2188 AIAMAEETT	A10EEQOQ A1670300
	411089	123172_1		454 AA713730 AA091294 AA584921 N86077 AW8367 195 AA514764 AA454562 A1082382 AA595822 AA5513				
80	411152	1234028_1	BE069	199 AW936012 AW877466 AW819782 AW935798 AW	835546 AW936042	BE069121 AW835625	AW877536 AW9	35885 BE069202
		_		0019 AW935937 BE160180 AW935946 BE069101 BE0				
	412537	1304_1	AL031	778 X59711 NM_002505 M59079 A1870439 A1494259 / 3 BE079412 BE079428 N90322 A1631202 A1141758 A1	444050 AA4050	OJ AA436132 BE1745 JB62076 AJ375230 AJ2	10 AA412691 AI4 08445 AW23576	3 AL044113 AA382556
			AW95	3 BEV/9412 BEV/9428 N9U322 ABS1202 AT141756 AT 3918 AA927051 AA889823 BE003094 AW390155 AW3	60805 AW360823 A	AW380810 AA425472	A1694282 AL044	14 A1684577 A1809865

	***	0 02/000443	AI478773 AI160445 AI674630 N69088 AW665529 N49278 AI129239 AI457890 AI621264 AW297152 AI268215 AA907787 AI286170 AI017982
			Al963541 Al469807 Al969353 BE552356 N66509 AA736741 AA382555 AW075811 AW292026
	412811	132943_1	H06382 AW957730 AA352014 R13591 AA121201 D60420 BE263253 BE047862 Z41952 Al424991 Al693507 Al863108 AA599060 Al091148
5			AA598689 R39887 AA813482 AW016452 H06383 R41807 Al364268 AA620528 Al241940 AW089149 AW090733 AW088875 Z38240
3	413690	1383256_1	AA121202 R17734 BE157489 BE157560
	414883	15024_1	AA926960 AA926959 W76521 W24270 W21526 AA037172 BE267636 H83186 AA469909 N86396 AA001348 BE535736 AA081745 BE566245
			AA082436 H72525 H77575 N49786 W80565 H78746 BE569085 W04339 R98127 T55938 BE279271 AW960304 T29812 AA476873 BE297387
10			AA292753 AA177048 NM_001826 X54941 BE314366 AA908783 AI719075 BE270172 BE269819 AA889955 AI204630 W25243 AI935150 AA872039 W72395 T99630 AI422691 H98460 N31428 BE255916 H03265 AI857576 AA776920 AA910644 AA459522 AA293140 AW514667
10			R75953 AW682395 AA662522 Al865147 Al423153 AW262230 AA584410 AA583187 AW024595 AW069734 Al828995 AA282997 A8676046
			AW613002 AA527373 AW972459 AIR31360 AA621337 AA100926 AA772418 AA594628 AI033892 W95096 AI034317 AA398727 AI085031
			N95210 Al459432 Al041437 AA932124 AA627684 AA935829 Al004827 Al423513 Al094597 H42079 R54703 Al630359 AA617681 AA978045
15			AA643280 W44561 Al991988 Al537692 Al090262 AA740817 Al312104 Al911822 AA416871 Al185409 AA129784 AA701623 Al075239 Al139549 AA633648 Al339996 Al336880 AA399239 Al078708 Al085351 Al362835 Al346618 Al146965 Al989380 Al348243 N92892 AA765850
15			A1494230 A1278887 AA982598 A1492600 W80435 AA001979 R97424 A1129015 N24127 AA157451 AA235549 AA459292 AA037114 AA129785
			AI494211 AW059601 AW886710 R92790 N59755 AI361128 AW589407 H47725 H97534 H48076 H48450 T99631 AW300758 H03431 R76789
			AA954344 H77576 R96823 AI457100 N92845 N49682 H42038 BE220699 BE220715 H99552 AA701624 N74173 R54704 H79520 H72923
20			H03266 BE261919 AA769633 AA480310 AA507454 AA910586 Al203723 AW104725 W25611 W25071 T88980 H03513 T77589 R99156 W95095 R97470 AA702275 T77551 AA911952 H82956 N83673 AA283872
	415989	156454_1	AI267700 AI720344 AA191424 AI023543 AI469633 AA172056 AW958465 AA172236 AW953397 AA355086
	417324	166714_1	AW265494 AA455904 AA195677 AW265432 AW991605 AA456370
	418574	1,7690_1	N28754 N28747 Al568146 Al979339 AA322671 AA322672 AW955043 Al990326 AA776406 Al016250 AA843678 AW451882 N23137 N23129 W70051 Al038748 AA831327 Al925845 AW945895
25	418712	1784125_1	742183 T31621 T97478
	419443	184788_1	D62703 AA242966 D79798
	419502	18535_1	AU076704 T74854 T74860 T72098 T73265 T73873 T69180 T74658 T58786 T60385 T73410 T68781 T67845 T67593 T73952 T67864 T60630
			T68367 T68401 T53959 T72360 T72099 T60377 T58961 T71712 T72821 T64738 T74645 T72037 T68688 T72063 T73258 T72826 T64242 T68220 T74673 T71800 T68355 T61227 T62738 T69317 T53850 T64692 T73768 T73962 T73382 T68914 T70975 T73400 T60631 T73277
30			T73203 T70498 T61409 T58925 NM, 000508 M94982 T68301 T73729 T69445 T60424 T67922 T67736 T68716 T67755 T74765 T73819 T58719
			T74766 T60477 T74863 T61109 T68329 T58850 T71857 T73425 T53736 T68607 T58898 T64309 T72031 T72079 T64306 T71908 T68107
			T71916 T73787 T56035 T64425 T71870 T60476 T61376 T67820 T71895 T41006 T69441 T68170 T74617 T71958 T69440 T61875 R06796 H48353 T71914 T53939 T64121 AA693996 T72525 T67779 T68078 AA011465 AA345378 AV654847 AV654272 AV656001 Al064740 T82897
			H9055 1/1914 150539 104121 A005398 11252 107779 100076 AVI 1405 AA394576 AV094047 AV094272 AV090011 A1004740 102697 N33594 AA344542 AW805054 A1207457 T61743 AA026737 H94389 AA382695 AA918409 T680444 S82092 T39959 A1017721 AA312395
35			AA312919 T40156 H66239 AV652989 H38728 R98521 AV655200 R95790 W03250 W00913 AA344136 AV660126 R97923 AA343596
			AW470774 AV651256 N54417 AA812862 AW182929 Al111192 H61463 H72060 AA344503 H38639 Al277511 AV661108 Al207625 T47810
			AA235252 T27853 T47778 R95746 H70620 AA701463 AW827166 R98475 C20925 AV657287 T71959 T71313 T73920 T73333 T61618 T69293 T69283 T73931 T72178 T72456 AV645639 AV653476 T72957 T72300 T68906 T71457 T70494 T72956 T70495 T68267 T74407 T85778
			10323 17331 17216 172431 XV95053 XV05374 17253 17250 17250 17453 17254 1
40			T70475 T64751 AA344441 AA343657 AA345732 AA344328 AI110639 AA344603 AF063513 T64696 T68516 T72223 T60507 T67633 R29500
	•		T72517 R02292 T60599 T69206 T70452 T74677 R29366 T61277 T74914 T60352 R29675 T74843 AV645792 AA344408 T69197 T72057
			T69368 T69358 T68258 AV650429 T73341 T61702 T74598 T40095 K02272 T40106 AA343045 AA341908 AA341907 AA342807 AA341964 T53747 T72042 T62764 A\064899 AA343060 T67832 T72440 T71770 T68091 T69108 T72449 T69167 T71289 T68251 AV654844 T64375
			A3345234 T67598 AA011414 T68036 H48262 Al207557 T68219 W86031 T69081 T64232 R93196 T62136 AV650539 H67459 T72978
45			AA344583 T60362 H58121 T95711 T72803 T68055 T71715 R29036 T72793 T69122 T64595 T62888 T69139 T68291 T64652 T67971 T46862
			AA693592 Al248502 R29454 T64764 T57001 T73052 T71429 T51176 T58866 AV655414 H90426 AA342489 T73666 T67848 T72512 T53835
	419936	189181_1	T67837 T73317 T74273 T69420 T68245 T74380 T67862 T74474 T56068 Al792788 BE142230 AA252019
	421582	2041_1	AI910275 X00474 X52003 X05030 NM_003225 AA314326 AA308400 AA506787 AA314825 AI571948 AA507595 AA614579 AA587613 R83818
50			AA568312 AA614409 AA307578 AI925552 AW950155 AI910083 M12075 BE074052 AW004668 AA578674 AA582084 BE074053 BE074126
			BE074140 AA514776 AA588034 BE074051 BE074068 AW009769 AW050690 AA858276 R55389 Al001051 AW050700 AW750216 AA614539 BE074045 Al307407 AW602303 BE073575 Al202532 AA524242 Al970839 Al909751 BE076078 Al909749 R55292
	422128	211994_1	BEU14043 AI3U14U1 AW6U23U3 BEU13313 AI2U2332 AA324242 AI31U633 AI3U3161 BEU16U16 AI3U3149 R35232 AW881145 AA490718 M85637 AA304575 T06067 AA331991
	423034	224122_1	AL119930 AA320696 AW752565
55	423816	23234_1	AL031985 AL137241 AI792386 AI733664 Al857654 Al049911
	424200 424999	236595_1 245835_1	AA337221 AA336756 AW966196 AW953120 R56325 AA349562
	426966	273896_1	A1993124 A1498691 AW771508 A1498457 A1768408 A1783624 A1383985 A1580267 D79813 AA393768
	426991	27415_1	AK001536 AA191092 AW510354 Al554256 AL353968 AA134266
60	427260	276598_1	AA663848 AA400100 AA401424
	428023	28589_2	AL038843 AA161338 BE268213 AA425597 N87306 AA092969 BE566038 AA247451 N47392 AI928802 AW182584 AW027872 AI819831 AI936994 W56258 AI653448 AI278611 AI283557 AI824306 AW338658 AW150899 AA687514 N47393 N29885 AA973469 AI038904 AI292064
			Al034339 AW674593 N72155 Al079733 Al038683 Al291616 AA491599 AA993675 AA837380 BE006554 BE006473 Al087090 T33044
<i>(</i>			AA652043 Al203503 AA583959 W35283 Al129926 Z41844 AW020925 AW575848 Al684603 AA493297 Al140689 Al277175 AA425444
65	400000	204204 4	AI932767 W02632 BE398786 R37261
	429220 429978	301384_1 31150_1	AW207206 AW341473 AA448195 Al951341 AA249027 AL038984 AK001993 AL080066 AV652725 BE566226 AA345557 AA315222 AA090585 AA375688 AA301092 AA298454 W05762
	420010	\$1130_1	AW607939 H51558 D83880 N84323 BE296821 AW947007 D61461 AW079261 AA329482 AW901780 AJ354442 AA772275 R31663 AJ354441
70			AI767525 H92431 AI916735 H93575 AI394255 AW014741 AI573090 C06195 AW612857 AW265195 AI339558 AI377532 AI308821 AI919424
70			AI589705 AW055215 AI336532 AI338051 AA806547 C76509 C00618 AW071172 AW769904 AA630381 AI678018 AI863985 D79662 BE221049 AW265018 AI589700 AW196655 N76573 AI370908 BE042393 N75017 AI698870 AW960115
	430439	31808_1	AW265018 AI589701 AW196655 N76573 AI570908 BE042393 N75017 AI696670 AW960115 AL133561 AL041090 AL117481 AL122069 AW439292 AI968826
	430935	325772_1	AW072916 Al184913 AA489195 AW466994 AW469044 N59350 Al819642 Al280239 Al220572 AA789302 Al473611 AW841126 D60937
75	431089	327825_1	BE041395 AA491826 AA621946 AA715980 AA666102
75	431322	331543_1	AW970622 AA503009 AA502998 AA502989 AA502805 T92188 AA221036 R87170 BE537068 BE544757 C18935 AW812058 T92565 AA227415 AA233942 AA223237 AA668403 AA601627 AW869639
	432407	34624_1	AAZZ1036 RB/170 BE537066 BE544737 C16935 AW612036 192565 AAZZ7415 AAZ33942 AAZZ3Z37 AA666403 AA601627 AW659539 BE661833 BE000620 AW961170 AW847519 AA308542 AW821833 AW945688 C04699 AA205504 AA377241 AW821667 AA055720
			AW817981 AW856488 AA155719 AA179928 T03007 AW754298 AA227407 AA113928 AA307904 C16859
90	434414	38585_1	AI798376 \$46400 AW811617 AW811616 W00557 BE142245 AW858232 AW861851 AW858362 AA232351 AA218567 AA055556 AW858231
80			AW857541 AW814172 H66214 AW814398 AF134164 AA243093 AA173345 AA199942 AA223384 AA227092 AA227080 T12379 AA092174 T61139 AA149776 AA699829 AW879188 AW813567 AW813538 AI267168 AA157718 AA157719 AA100472 AA100774 AA130756 AA157705
			AA157730 AA157715 AA053524 AW849581 AW854566 C05254 AW88286 T92637 AW812621 AA206583 AA209204 BE156909 AA226824
			AI829309 AW991957 N66951 AA527374 H66215 AA045564 AI694265 H60808 AA149726 AW195620 BE081333 BE073424 AW817662
85	436600	40264 2	AW817705 AW817703 AW817659 BE081531 H59570 A Academy Altagess Becompte
0.5	436608	42361_3	AA628980 AI126603 BE504035

	***	D 02/0964	2 DCT/IIS02/12/17	
	438091	O 02/0864 44964_1	3 PCT/US02/12476 AW373062 T55662 AI299190 BE174210 AW579001 H01811 W40186 R67100 AI923886 AW952164 AA628440 AW888607 AW898616 AA709126 AW898628 AW888544 AA947932 AW888625 AW898622 AI276125 AI185720 AW510698 AA987230 T52522 BE487708 AW243400 AW043642 AI288245 AI186932 D52654 D55017 D52715 D52477 D53933 D54679 AI298739 AI146984 AI922204 N98343 BE174213 AA84557 AI813854 AI214518 AI635262 AI139455 AI707807 AI698085 AW884528 AI024768 AI004723 AW087420 AI565133 N94964 AI268939) /1
5			AW513280 Al061126 Al435818 Al859106 Al360506 Al024767 AA513019 AA757598 X56196 AA902959 Al334784 Al860794 AA010207 AW890091 AW513771 Al951391 Al337671 T52499 AA890205 Al640908 H75966 AA463487 AA358688 Al961767 Al866295 AA780994 Al985913 BE174196 AA029094 AW592159 T55581 N79072 Al611201 AA910812 Al220713 AW149306 Al758412 AA045713 R79750 N76096	ı
10	439000 439285	467716_1 47065_1	AW979121 AA847986 AA829098 AL133916 N79113 AF086101 N76721 AW950828 AA364013 AW955684 AI346341 AI867454 N54784 AI655270 AI421279 AW014882 AA775552 N62351 N59253 AA626243 AI341407 BE175639 AA456968 AI358918 AA457077	
	439780 441128 443068	47673_1 51021_2 558874_1	AL109688 R23665 R26578 AA570256 AW014761 AA573721 AI473237 AI022165 AA554071 AA127551 N90525 AW973623 AA447991 AA243852 BE328850 AI148171 AI359627 AI005068 AI356567 AA232991 AW016855 AA906902 AA233101 AA127550 BE512923 AI188710 AI032142 AW078833 N30308 AW675632 AI219028 AI341201 N22181 H95390	
15	443947 447636	586160_1 7301_1	W24187 W24194 R17789 Y10043 NM_005342 L05085 AL034450 BE614226 AW749053 AA379173 AA248230 BE514634 AA334622 R70656 AA367593 AA214649 AA369318 AW957081 R05760 AA039903 AI886597 AW630122 AA906264 AA041527 R01145 AI088688 BE463637 AA398795 AI354883 AJ768938 AI569996 AI452952 AI168582 AI189869 AI086670 AW262560 AW613854 AA862839 AA435840 AA670197 AI024032 AI990659 AI990089 N81095 AA847919 AW960150 AA211075 AA044704 AA367594 AW582587 AW858854 AW818630 AW818281 AW818433 AW5825	595
20	448993	79225_1	AA096002 N83992 AI471630 BE540637 BE265481 AW407710 BE513882 BE546739 AA053597 BE140503 BE218514 AW956702 AI656234 AI636283 AI567265 AW340858 BE207794 AA053085 R69173 AA292343 AA454908 AA293504 AI659741 AI927478 AA399460 AI760441 AA346416 BE047245 AA730380 AA394063 AA454833 AI982791 AI567270 AI813332 AI767858 AA427705 D20284 AI221458 BE048537 AI263048 AA346417 AA911497 BE537702	
25	449305 451105 451320	804424_1 859083_1 86576_1	Al638293 AW813561 Al761324 AW880941 AW880937 AW118072 Al631982 T15734 AA224195 Al701458 W20198 F26326 AA890570 N90552 AW071907 Al671352 Al375892 T03517 R88265 Al124088 AA224388 Al084316 Al354686 T33652 Al140719 Al720211 T03490 Al372637 T15415 AW205836 AA630384 T03515 T33230 AA017131 AA443303 T33623 Al222556 T33511 T33785 Al419606 D55612	
30	451807 .	8865_1	W52854 AL117600 BE208116 BE208432 BE206239 BE082291 AW953423 AA351619 BE180648 BE140550 W60080 AA865478 N90291 AW450652 AW449519 AA993634 AI806539 AA351618 AW449522 AI827626 AA904788 AA380381 AA886045 AA774409 BE003229 Z41756 AL133819 AA488118 AA383064 AI476447 T09430 AI673758 AA524895 AI581345 AI300820 AW498812 AA256162 AI559724 AI685732	
35	452410	9163_1	AA602400 AA905453 Al204595 AW166541 AA157456 AA156269 AA383652 AA431072 AW592707 Al435410 AW272464 Al215594 AA62274 R74039 N35031 Al804128 AW513621 AA868351 Al026826 Al493388 AA614641 W81604 Al567080 Al214351 AA730140 Al125754 Al200813 Al269603 Al565082 Al807095 Al476629 AA505909 Al368449 Al686077 Al582930 AW085038 AA757863 AA730154 Al767072 AA468316	7
55	454241 455175	1067807_1 1257335_1	A1734130 A1734138 AA426284 AA433997 A1741241 AW043563 A1732741 A1732734 AA437369 AA425820 AA664048 R74130 BE144666 BE184942 AW238414 BE184946 AW993247 AW861464	
40	456237 458098	168730_1 47395_1	AA203682 R11958 BE550224 AA832519 N45402 AW885857 N29245 BE465409 W07677 AW970089 Al299731 AA482971 BE503548 H18151 W79223 AF08639 AA461301 W74510 R34182 Al090689 N46003 BE071550 R28075 AW134982 Al240204 Al138906 AW026179 Al572316 BE466182 Al206395 Al276154 Al273269 Al422817 Al371014 Al421274 Al188525 AA939164 BE549810 AW137865 Al694996 BE503841 AA459718 BE327407 BE467534 BE218421 BE467767 AA989054 BE467063 Al797130 BE327781	3
45	TABLE 9C			
50	Pkey: Ref: Strand: Nt_position	Sequence se sequence of Indicates DN	r corresponding to an Eos probeset rce. The 7 digit numbers in this column are Genbank Identifier (GI) numbers. "Dunham I. et al." refers to the publication entitled "The DNA uman chromosome 22." Dunham I. et al., Nature (1999) 402:489-495. strand from which exons were predicted. solide positions of predicted exons.	
55	Pkey 400512	Ref S	and NL position us 1439-1615	
33	400512 400517 400560 400664 400665	9796686 M 9843598 P 8118496 P	us 49996-50346 94182-94323,97056-97243,101095-101236,102824-103005 13558-13721,13942-14090,14554-14679	
60	400666 400749 400763 401027	8118496 P 7331445 M 8131616 M		
65	401027 401093 401203 401212 401411	8516137 M 9743387 M 9858408 P	us 22335-23166 us 172961-173056,173868-173928	
70	401435 401464 401714 401747	8217934 M 6682291 M 6715702 P	us 54508-55233 us 170688-170834	
75	401760 401780 401781	9929699 P 7249190 M	131932,132451-132575,133580-134011	
, 5	401785 401797 401961	7249190 M 6730720 P 4581193 M	165776-165996, 166189-166314, 166408-166569, 167112-167268, 167387-167469, 168634-168942 6973-7118 us 124054-124209	
80	401985 401994 402075 402260	2580474 P 4153858 M 8117407 P 3399665 M	61542-61750 us 42904-43124,43211-43336,44607-44763,45199-45281,46337-46732 5 121907-122035,122804-122921,124019-124161,124455-124610,125672-126076 us 113765-113910,115653-115765,116808-116940	
85	402265 402297 402408	3287673 P 6598824 P 9796239 M		

	WO 02/086443				PCT/US02/12476
	402420	9796339	Plus	129750-129919	
	402674	8077108	Minus	39290-39502	
	402802	3287156	Minus	53242-53432	
	402994	2996643	Minus	4727-4969	
5	403137	9211494	Minus	92349-92572,92958-93084,93579-93712,93949-94072,94591-94748,95214-95337	
_	403306	8099945	Plus	127100-127251	
	403329	8516120	Plus	96450-96598	
	403381	9438267	Minus	26009-26178	
	403478	9958258	Plus	116458-116564	
10	403485	9966528	Plus	2888-3001,3198-3532,3655-4117	
	403627	8569879	Minus	23868-24342	
	403715	7239669	Plus	85128-85292	
	404044	9558573	Minus	225757-225939	
	404076	9931752	Minus	3848-3967	
15	404101	8076925	Minus	125742-125997	
	404140	9843520	Plus	37781-38147	
	404165	9926489	Minus	69025-69128	
	404185	4572584	Minus	129171-129327	
	404210	5006246	Plus	169926-170121	
20	404253	9367202	Minus	55675-56055	
	404287	2326514	Plus	53134-53281	
	404298	9944263	Minus	73591-73723	
	404347	9838195	Plus	74493-74829	
	404440	7528051	Plus	80430-81581	
25	404721	9856648	Minus	173763-174294	
	404794	4826439	Plus	101619-101898	
	404854	7143420	Plus	14260-14537	
	404877	1519284	Plus	1095-2107	
	404927	7342002	Plus	68690-69563	
30	404996	6007890	Plus	37999-38145,38652-38998,39727-39872,40557-40674,42351-42450	
	405449	7622497	Plus	42236-42570	
	405568	6006906	Plus	35912-36065	
	405572	3800891	Plus	85230-85938	
	405646	4914350	Plus	741-969	
35	405676	4557087	Plus	73195-73917	
	405770	2735037	Plus	61057-62075	
	405932	7767812	Minus	123525-123713	
	406137	9166422	Minus	30487-31058	
	406360	9256107	Minus	7513-7673	
40	408399	9256288	Minus	63448-63554	
	406467	9795551	Plus	182212-182958	

TABLE 10A: Potential Therapeutic, Diagnostic and Prognostic targets for Therapy of Lung Cancer and Non-mailgnant Lung Disease
Table 2A shows about 307 genes up-regulated in non-mailgnant lung disease relative to lung tumors and normal body tissues and/or down-regulated in lung tumors relative to
normal lung and non-mailgnant lung disease. These genes were selected from about 59680 probesets on the Eos/Affymetrix Hu03 Genechip array. 45

Table 108 show the accession numbers for those Pkey's lacking UnigenetD's for table 10A. For each probeset we have listed the gene cluster number from which the oilgonucleotides were designed. Gene clusters were compiled using sequences derived from Genbank ESTs and mRNAs. These sequences were clustered based on sequence similarity using Clustering and Alignment Tools (DoubleTwist, Oakland California). The Genbank accession numbers for sequences comprising each cluster are listed in the 50

Table 10C show the genomic positioning for those Pkey's lacking Unigene ID's and accession numbers in table 10A. For each predicted exon, we have listed the genomic sequence source used for prediction. Nucleotide locations of each predicted exon are also listed.

Unique Eos probeset identifier number Exemplar Accession number, Genbank accession number Pkey: UnigenelD: Unigene number Unigene Title:

R1:

55

60

Unigene title
Average of lung tumors (including squamous cell carcinomas, adenocarcinomas, small cell carcinomas, granulomatous and carcinold tumors) divided by the
average of normal lung samples
Average of normalignant lung disease samples (including bronchitis, emphysema, fibrosis, atelectasis, asthma) divided by the average of normal lung samples R2:

65	Pkey	ExAccn	UnigenelD	Unigene Tille	R1	R2
	404394		-	ENSP00000241075:TRRAP PROTEIN.	0.79	3.10
	404916			Tarpet Exon	1.00	159.00
	405257			Target Exon	1.00	422.00
	407228	M25079	Hs.155376	hemoglobin, beta	0.47	2.33
70	407568	AA740964	Hs.62699	ESTs	1.00	123.00
	408562	AI436323	Hs.31141	Homo sapiens mRNA for KIAA1568 protein,	1.00	230.00
	409031	AA376836	Hs.76728	ESTs	1.00	128.00
	410434	AF051152	Hs.63668	toll-like receptor 2	39.65	149.00
	410467	AF102546	Hs.63931	dachshund (Drosophila) homolog	1.00	109.00
75	410808	T40326	Hs.167793	ESTs	1.14	13.14
	412351	AL135960	Hs.73828	T-cell acute lymphocytic leukemia 1	0.37	2.27
	412372	R65998	Hs.285243	hypothetical protein FLJ22029	1.00	173.00
	413795	AL040178	Hs.142003	ESTs	0.10	11.90
	414154	AW205314	Hs.323060	ESTs	0.62	2.09
80	414214	D49958	Hs.75819	glycoprotein M6A	0.03	4.55
	414998	NM_002543	Hs.77729	oxidised low density lipoprotein (lectin	0.64	2.97
	415122	D60708	Hs.22245	ESTs	0.07	8.97
	415765	NM 005424	Hs.78824	tyrosine kinase with immunoglobulin and	0.67	1.65
	415775	H00747	Hs.29792	ESTs, Weakly similar to 138022 hypotheti	0.29	2.64
85	415910	U20350	Hs.78913	chemokine (C-X3-C) receptor 1	1.00	145.00

		O 02/086				
	416319	AI815601	Hs.79197	CD83 antigen (activated B lymphocytes, i	15.32 0.64	237.00 4.00
	416402 417355	NM_000715 D13168	Hs.1012 Hs.82002	complement component 4-binding protein, endothelin receptor type B	0.04	3.90
	417421	AL138201	Hs.82120	nuclear receptor subfamily 4, group A, m	36.30	357.00
5	417511	AL049176	Hs.82223	chordin-like	1.00	179.00
	418489	U76421	Hs.85302	adenosine deaminase, RNA-specific, B1 (h	0.02	6.00
	418726	BE241812 H83265	Hs.87860 Hs.8881	protein tyrosine phosphalase, non-recept ESTs, Weakly similar to S41044 chromosom	1.00 0.44	113.00 1.90
	418741 418883	BE387036	Hs.1211	acid phosphatase 5, tartrate resistant	0.96	2.04
10	419086	NM_000216	Hs.89591	Kalimann syndrome 1 sequence	0.62	2.74
	419150	T29618	Hs.89640	TEK tyrosine kinase, endothelial (venous	0.03	6.90
	419235	AW470411	Hs.288433	neurotrimin hypothetical protein FLJ21276	1.48 37.55	5.13 336.00
	419407 420556	AW410377 AA278300	Hs.41502 Hs.124292	Homo saplens cDNA: FLJ23123 fis, clone L	0.80	3.65
15	420656	AA279098	Hs.187636	ESTs	1.65	8.07
	420729	AW964897	Hs.290825	ESTs	2.99	25.82
	421177	AW070211	Hs.102415	Homo sapiens mRNA; cDNA DKFZp586N0121 (f	0.46 1.00	1.95 156.00
	422060 422426	R20893 W79117	Hs.325823 Hs.58559	ESTs, Moderately similar to ALU5_HUMAN A ESTs	0.03	7.44
20	422652	AW967969	Hs.118958	syntaxin 11	0.14	3.62
	423099	NM_002837	Hs.123641	protein tyrosine phosphatase, receptor t	0.01	3.16
	424433	H04607	Hs.9218	ESTs ESTs	0.75 1.00	141.75 167.00
	424585 424711	AA484840 NM_005795	Hs.131987 Hs.152175	calcitonin receptor-like	0.43	3.01
25	424973	X92521	Hs.154057	matrix metalloproteinase 19	0.37	19.45
	425023	AW956889	Hs.154210	endothelial differentiation, sphingolipi	0.14	3.35
	425664	AJ006276	Hs.159003 Hs.165950	transient receptor potential channel 6 fibroblast growth factor receptor 4	1.00 0.68	94.00 1.42
	425998 426657	AU076629 NM_015865	Hs.171731	solute carrier family 14 (urea transport	0.03	3.74
30	426753	T89832	Hs.170278	ESTs	1.00	141.00
	427558	D49493	Hs.2171	growth differentiation factor 10	1.00	117.00
	427983	M17706 AK002121	Hs.2233 Hs.184465	colony stimulating factor 3 (granulocyte hypothetical protein FLJ11259	0.75 0.76	2.20 2.25
	428467 428927	AA441837	Hs.90250	ESTs	0.01	3.62
35	429496	AA453800	Hs.192793	ESTs	1.00	138.00
	430468	NM_004673	Hs.241519	angiopoletin-like 1	1.00	132.00
	431385 431728	BE178536 NM_007351	Hs.11090 Hs.268107	membrane-spanning 4-domains, subfamily A multimerin	1.00 1.00	167.00 157.00
	431848	Al378857	Hs.126758	ESTs, Highly similar to AF175283 1 zinc	0.34	2.24
40	432128	AA127221	Hs.117037	ESTs	0.00	1.15
	432519	AJ221311	Hs.130704	ESTs, Weakly similar to BCHUIA S-100 pro	0.01 1.00	2.06 267.00
	433043 433803	W57554 Al823593	Hs.125019 Hs.27688	lymphold nuclear protein (LAF-4) mRNA ESTs	1.00	105.00
	434730	AA644669	Hs.193042	ESTs	1.05	3.15
45	435472	AW972330	Hs.283022	triggering receptor expressed on myelold	0.83	1.94
	436532	AA721522	Un 477042	gb:nv54h12.r1 NCI_CGAP_Ew1 Homo saplens ESTs	1.00 1.00	218.00 133.00
	437119 437140	Al379921 AA312799	Hs.177043 Hs.283689	activator of CREM in testis	0.67	122.67
~ ^	437211	AA382207	Hs.5509	ecotropic viral integration site 2B	1.00	142.00
50	437960	AI669586	Hs.222194	ESTs	1.00	147.00
	438202 438873	AW169287 Al302471	Hs.22588 Hs.124292	ESTs Homo saplens cDNA: FLJ23123 fis, clone L	1.00 0.71	141.00 3.66
	438875	AA827640	Hs.189059	ESTs	23.32	370.00
~ ~	441048	AA913488	Hs.192102	ESTs	0.77	8.50
55	441188	AW292830	Hs.255609	ESTs	3.43 1.00	16.36 167.00
	441499 444513	AW298235 AL120214	Hs.101689 Hs.7117	ESTs glutamate receptor, lonotropic, AMPA 1	1.00	151.00
	444527	NM_005408	Hs.11383	small inducible cytokine subfamily A (Cy	46.47	153.00
~	444561	NM_004469	Hs.11392	c-fos induced growth factor (vascular en	0.01	3.08
60	445279	R41900	Hs.22245	ESTS	0.60 0.18	141.00 2.39
	446017 446984	N98238 AB020722	Hs.55185 Hs.16714	ES1s Rho guanine exchange factor (GEF) 15	0.10	2.16
	446998	N99013	Hs.16762	Homo seplens mRNA; cDNA DKFZp564B2062 (f	0.01	2.53
<i>CE</i>	447357	Al375922	Hs.159367	ESTs	0.46	2.64
65	448108 448253	A1800470 H25899	Hs.171941 Hs.201591	ESTs ESTs	18.05 1.00	296.00 141.00
	449275	AW450848	Hs.205457	perlaxin	0.56	1.38
	450400	Al694722	Hs.279744	ESTs	0.88	4.33
70	450696	AI654223	Hs.16026	hypothetical protein FLJ23191	0.52 0.79	2.08 2.01
70	450726 451497	AW204600 H83294	Hs.250505 Hs.284122	retinoic acid receptor, alpha Wnt inhibitory factor-1	0.75	2.03
	451533	NM_004657	Hs.26530	serum deprivation response (phosphatidyl	0.13	2.25
	453636	R67837	Hs.169872	ESTs	1.00	116.00
75	458332 459580	AI000341 AA022888	Hs.220491 Hs.176065	ESTs ESTs	1.00 0.20	192.00 2.98
75	400269	771022000	115.170000	Eos Control	0.40	2.40
	403421			NM_016369*:Homo sapiens claudin 18 (CLDN	0.53	1.77
	407570	Z19002	Hs.37096	zinc finger protein 145 (Kruppel-like, e	0.01	3.18
80	412295 414517	AW088826 M24461	Hs.117176 Hs.78305	poly(A)-binding protein, nuclear 1 surfactant, pulmonary-associated protein	0.56 0.64	1.74 1.50
00	417204	N81037	Hs.1074	surfactant, pulmonary-associated protein	0.33	1.16
	418307	U70867	Hs.83974	solute carrier family 21 (prostaglandin	0.53	1.55
	418935	T28499	Hs.89485	carbonic anhydrase IV	0.20 0.78	1.28 1.90
85	421502 421798	AF111856 N74880	Hs.105039 Hs.29877	solute carrier family 34 (sodium phospha N-acylsphingosine amidohydrolase (acid c	0.78	1.54

	w	O 02/086	443			
	423354	AB011130	Hs.127436	calcium channel, voltage-dependent, alph	0.59	1.55
	423738	AB002134	Hs.132195	airway trypsin-like protease	10.14 0.35	51.00 1.62
	425211 425438	M18667 T62216	Hs.1867 Hs.270840	progastricsin (pepsinogen C) ESTs	0.33	9.45
5	426828	NM_000020	Hs.172670	activin A receptor type II-like 1	0.03	1.71
•	427019	AA001732	Hs.173233	hypothetical protein FLJ10970	0.01	1.49
	428043	T92248	Hs.2240	uleroglobin	0.42	1.26 2.43
	430280 431433	AA381258 X65018	Hs.237868 Hs.253495	interleukin 7 receptor surfactant, pulmonary-associated protein	0.46 0.57	1.59
10	431723	AW058350	Hs.16762	Homo sapiens mRNA; cDNA DKFZp564B2062 (f	0.29	1.80
	432985	T92363	Hs.178703	ESTs	0.32	2.27
	441835	AB036432	Hs.184	advanced glycosylation end product-speci	0.31	1.51
	442275	AW449467	Hs.54795	ESTs ESTs	0.55 0.00	1.78 3.02
15	443709 444325	AI082692 AW152618	Hs.134662 Hs.16757	ESTs	0.32	2.49
15	450954	AJ904740	Hs.25691	receptor (calcitonin) activity modifying	0.46	1.74
	451558	NM_001089	Hs.26630	ATP-binding cassette, sub-family A (ABC1	0.52	1.87
	453310	X70697	Hs.553	solute carrier family 6 (neurotransmitte	0.00 0.01	3.30 2.31
20	456855 444342	AF035528 NM_014398	Hs.153863 Hs.10887	MAD (mothers against decapentaplegic, Dr similar to lysosome-associated membrane	0.66	2.20
20	400754	14111_014000	113.10031	Target Exon	1.00	297.00
	401045			C11001883*:gij6753278 ref NP_033938.1 c	1.00	109.00
	401083			NM_016582*:Homo sapiens peptide transpor	0.89 1.45	1.39 4,47
25	402474 402808			NM_004079:Homo sapiens cathepsin S (CTSS ENSP00000235229:SEMB.	1.00	1.87
23	403021			C21000030:gi 9955960 ref NP_063957.1 AT	1.00	149.00
	403438			NM_031419*:Homo sapiens molecule possess	1.06	2.96
	403687			NM_007037*:Homo sapiens a disintegrin-li	0.04	4.89 225.00
30	403764			NM_005463:Homo sapiens heterogeneous nuc NM_019111*:Homo sapiens major histocompa	1.00 0.97	1.93
30	404277 404288			NM_002944*:Homo sapiens v-ros avian UR2	1.00	68.00
	404518	Al815601		CD83 antigen (activated B lymphocytes, i	0.02	1.83
	405106			C11001637*:gij5032241[ref[NP_005732.1] z	1.00	235.00
35	405381			Target Exon	1.00 1.37	93.00 6.02
33	406387 406646	M33600		Target Exon mejor histocompatibility complex, class	0.86	2.48
	408714	Al219304	Hs.266959	hemoglobin, gamma G	0.01	3.19
	406753	AA505665	Hs.217493	annexin A2	1.00	147.00
40	406973	M34996	Hs.198253	major histocompatibility complex, class	1.03 1.00	2.04 64.00
40	407248 407510	U82275 U96191	Hs.94498	teukocyte immunoglobulin-like receptor, gb:Human trophoblast hypoxia-regulated f	1.00	90.00
	407731	NM_000068	Hs.38069	complement component 8, beta polypeptide	1.00	67.00
	407830	NM_001086	Hs.587	arylacetamide deacetylase (esterase)	1.00	102.00
45	408045	AW138959	Hs.245123	ESTs	1.00 1.00	70.00 112.00
43	408074 408374	R20723 AW025430	Hs.155591	ESTs forkhead box F1	0.07	10.17
	409064	AA062954	Hs.141883	ESTs	0.39	2.31
	409083	AF050083	Hs.673	interleukin 12A (natural killer cell sti	1.00	95.00
50	409153	W03754	Hs.50813	hypothetical protein FLJ20022 cytochrome P450, subfamily IVB, polypept	0.01 0.01	4.55 3.72
50	409203 409238	AA780473 AL049990	Hs.687 Hs.51515	Homo sapiens mRNA; cDNA DKFZp564G112 (fr	1.00	79.00
	409389	AB007979	Hs.301281	Homo saplens mRNA, chromosome 1 specific	0.14	27.35
	409718	D86640	Hs.56045	src homology three (SH3) and cysteine ri	1.00	113.00
55	410798 411020	BE178622 NM_006770	Hs.16291 Hs.67726	gb:PM3-HT0605-270200-001-a02 HT0605 Homo macrophage receptor with collagenous str	0.64 0.55	2.47 2.40
55	411667	8E160198	115.07720	gb:QV1-HT0413-010200-059-h03 HT0413 Romo	1.00	111.00
	412000	AW576555	Hs.15780	ATP-binding cassette, sub-family A (ABC1	1.00	95.00
	412358	BE047490	Hs.24172	ESTs	1.00	87.00
60	412420 412564	AL035668 X83703	Hs.73853 Hs.31432	bone morphogenetic protein 2 cardiac ankyrin repeat protein	1.43 0.02	8.07 3.07
oo	412869	AA290712	Hs.82407	CXC chemokine ligand 16	0.93	1.72
	412870	N22788	Hs.82407	CXC chemokine ligand 16	0.97	1.51
	413529	U11874	Hs.846	interleukin 8 receptor, beta	0.02 0.65	2.42 1.50
65	413533 413689	BE146973 BE157286	Hs.20631	gb:QV4-HT0222-011199-019-e05 HT0222 Homo zinc finger protein, subfamily 1A, 5 (Pe	20.87	232.00
UJ	413724	AA131466	Hs.23767	hypothetical protein FLJ12666	1.00	80.00
•	413800	Al129238	Hs.192235	ESTs	1.00	85.00
	413802	AW964490	Hs.32241	ESTs, Weakly similar to S65657 alpha-1C-	1.00 0.02	213.00 3.93
70	413829 414376	NM_001872 BE393856	Hs.75572 Hs.66915	carboxypeptidase B2 (plasma) ESTs, Wealdy similar to 16.7Kd protein [1.00	115.00
,,	414577	AI056548	Hs.72116	hypothetical protein FLJ20992 similar to	0.49	1.94
	414700	H63202	Hs.38163	ESTs	0.03	3.75
	415078	AA311223	Hs.283091	found in Inflammatory zone 3	0.86	1.95 120.00
75	415120 415323	N64464 BE269352	Hs.34950 Hs.949	ESTs neutrophil cytosolic factor 2 (65kD, chr	1.00 0.60	2.48
, 5	415335	AA847758	Hs.111030	ESTs	1.00	95.00
	415582	W92445	Hs.165195	Homo sapiens cDNA FLJ14237 fis, clone NT	1.00	136.00
	416030	H15261	Hs.21948	ESTs PortCole 42 guardine exchange factor (GEF)	0.02 1.00	8.07 73.00
80	416427 416464	BE244050 NM_000132	Hs.79307 Hs.79345	Rac/Cdc42 guanine exchange factor (GEF) coagulation factor VIII, procoagulant co	0.70	3.36
	416585	X54162	Hs.79386	leiomodin 1 (smooth muscle)	0.06	6.56
	416847	L43821	Hs.80261	enhancer of filamentation 1 (cas-like do	0.70	3.66
	417148	AA359896	Hs.293885	hypothetical protein FLJ14902 tryptophanyl-IRNA synthetase	1.00 0.85	114.00 1.30
85	417370 417673	T28651 T87281	Hs.82030 Hs.16355	ESTs	0.15	15.54
Ų,	4.1013					

		O 02/086				
	418067	Al127958	Hs.83393	cystatin E/M	0.81 1.00	1.74 99.00
	418296 418643	C01566 J03798	Hs.86671 Hs.86948	ESTs small nuclear ribonucleoprotein D1 polyp	1.00	60.00
_	418832	X04011	Hs.88974	cytochrome b-245, beta polypeptide (chro	2.40	14.74
5	418945	BE246762	Hs.89499	arachidonate 5-lipoxygenase	0.67	3.16
	419261	X07876 U08989	Hs.89791 Hs.91139	wingless-type MMTV integration site fami solute carrier family 1 (neuronal/epithe	1.00 1.00	73.00 192.00
	419564 419574	AK001989	Hs.91165	hypothetical protein	1.00	94.00
	419968	X04430	Hs.93913	interleukin 6 (interferon, beta 2)	61.16	500.00
10	420256	U84722	Hs.76206	cadherin 5, type 2, VE-cadherin (vascula	0.52	1.70 172.00
	420285 420577	AA258124 AA278436	Hs.293878 Hs.186649	ESTs, Moderately similar to ZN91_HUMAN Z ESTs	1.00 1.00	97.00
	421262	AA286746	Hs.9343	Homo saplens cDNA FLJ14265 fis, clone PL	1.00	64.00
	421445	AA913059	Hs.104433	Homo sapiens, clone IMAGE:4054868, mRNA	0.88	1.51
15	421470	R27496	Hs.1378	annexin A3	0.05 1.00	11.26 73.00
	421478 421563	Al683243 NM_006433	Hs.97258 Hs.105806	ESTs, Moderately similar to S29539 ribos granulysin	0.82	2.42
	421566	NM_000399	Hs.1395	early growth response 2 (Krox-20 (Drosop	5.50	31.57
20	421855	F06504	Hs.27384	ESTs, Moderately similar to ALU4_HUMAN A	1.00	129.00
20	421913	A1934365	Hs.109439	osteoglycin (osteoinductive factor, mime ESTs, Moderately similar to AF161511 1 H	1.00 0.60	101.00 63.60
	421952 422232	AA300900 D43945	Hs.98849 Hs.113274	transcription factor EC	1.00	148.00
	422386	AF105374	Hs.115830	heparan sulfate (glucosamine) 3-O-sulfot	1.40	3.98
25	423168	R34385	Hs.124940	GTP-binding protein	0.34	3.59
25	423196	AK001866	Hs.125139	hypothetical protein FLJ11004	0.55 0.09	2.00 2.13
	423387 423424	AJ012074 AF150241	Hs.128433	vasoactive intestinal peptide receptor 1 prostaglandin D2 synthase, hematopoietic	1.00	141.00
	423456	AL110151	Hs.128797	DKFZP586D0824 protein	1.00	66.00
20	423696	Z92546		Sushi domain (SCR repeat) containing	0.73	1.27
30	424027	AW337575	Hs.201591 Hs.143131	ESTs glycoprotein A33 (transmembrane)	0.54 0.77	2.58 2.47
	424212 425087	NM_005814 R62424	Hs.126059	ESTs	1.00	74.00
	425175	AF020202	Hs.155001	UNC13 (C. elegans)-like	0.85	1.96
25	425771	BE561776	Hs.159494	Bruton agammaglobulinemia tyrosine kinas	1.18	2.56
35	426486 427507	BE178285 AF240467	Hs.170056 Hs.179152	Homo sapiens mRNA; cDNA DKFZp586B0220 (f toll-like receptor 7	1.00 1.00	76.00 63.00
	427618	NM_000760	Hs.2175	colony stimulating factor 3 receptor (gr	0.60	2.19
	427732	NM_002980	Hs.2199	secretin receptor	0.97	1.42
40	427952	AA765368	Hs.293941	ESTs, Moderately similar to A53959 throm	1.00	105.00
40	428709 428769	BE268717 AW207175	Hs.104916 Hs.106771	hypothetical protein FLJ21940 ESTs	1.00 0.09	80.00 2.55
	428780	Al478578	Hs.50636	ESTs	1.00	98.00
	428833	A1928355	Hs.185805	ESTs	1.00	113.00
45	429657	D13626	Hs.2465	KIAA0001 gene product; putative G-protei	1.00	52.00
43	430212 430226	AA469153 BE245562	Hs.2551	gb:nc67f04.s1 NCI_CGAP_Pr1 Homo saplens adrenergic, beta-2-, receptor, surface	1.00 0.11	132.00 15.60
	430376	AW292053	Hs.12532	chromosome 1 open reading frame 21	1.00	103.00
	430414	AW365665	Hs.120388	ESTs	0.50	6.96
50	430656	AA482900	Hs.162080	ESTs	1.00 1.00	70.00 90.00
50	430843 430998	A1734149 AF128847	Hs.119514 Hs.204038	ESTs indolethylamine N-methyltransferase	0.29	1.84
	431217	NM_013427	Hs.250830	Rho GTPase activating protein 6	1.00	79.00
	431921	N46466	Hs.58879	ESTs	0.91	1.67
55	432176 432203	AW090386 AA305746	Hs,112278 Hs,49	arrestin, beta 1 macrophage scavenger receptor 1	0.66 1.00	2.63 76.00
55	432231	AA339977	Hs.274127	CLST 11240 protein	0.46	1.46
	432485	N90866	Hs.276770	CDW52 antigen (CAMPATH-1 antigen)	0.79	2.25
	432522	D11466	Hs.51	phosphatidylinositol glycan, class A (pa	1.93	4.83
60	432596 432850	AJ224741 X87723	Hs.278461 Hs.3110	matrilin 3 angiotensin receptor 2	0.04 1.00	5.7 9 167.00
00	433138	AB029496	Hs.59729	semaphorin sem2	0.04	9.16
	433563	A1732637	Hs.277901	ESTs	1.00	91.00
	433588	AI056872	Hs.133386	ESTs ESTs	120.16 0.60	315.00 1.84
65	434445 435496	Al349306 AW840171	Hs.11782 Hs.265398	ESTs. Weakly similar to transformation-r	1.00	128.00
••	435974	U29690	Hs.37744	Homo sapiens beta-1 adrenergic receptor	1.00	108.00
	436061	AI248584	Hs.190745	Homo saplens cDNA: FLJ21326 fis, clone C	1.00	91.00
	437157	BE048860 T27503	Hs.120655 Hs.15929	ESTs hypothetical protein FLJ12910	1.00 1.00	87.00 105.00
70	437207 437311	AA370041	Hs.9456	SWI/SNF related, matrix associated, acti	1.00	71.00
	437439	H29796	Hs.269622	ESTs	1.00	115.00
	438199	AW016531	Hs.122147	ESTs	1.00	80.00
	439551	W72062 AJ131245	Hs.11112 Hs.7239	ESTs SEC24 (S. cerevisiae) related gene famil	0.30 1.00	3.10 77.00
75	440515 440887	AJ 13 1245 Al 799488	Hs.135905	ESTs	1.00	85.00
	441025	AA913880	Hs.176379	ESTs	1.00	82.00
	441384	AA447849	Hs.288660	Homo sapiens cDNA: FLJ22182 fis, clone H	0.79	1.89 75.00
	441735 442200	AI738675 AW590572	Hs.127346 Hs.235768	ESTs ESTs	1.00 0.78	75.00 5.83
80	442832	AW206560	Hs.253569	ESTS	0.03	10.88
	442957	Al949952	Hs.49397	ESTs	1.00	70.00
	443282	T47764	Hs.132917	ESTs	1.00	197.00
	443547 443951	AW271273 F13272	Hs.23767 Hs.111334	hypothetical protein FLJ12666 ferritin, light polypeptide	1.00 0.55	253.00 2.09
85	444330	AI597655	Hs.49265	ESTs	1.00	90.00
		-				

WO 02/086443 PCT/US02/12476 1.00 84.00 444515 AW20490B Hs.169979 **ESTs** 0.02 445769 Hs.23666 4.38 AI741471 Homo saplens clone 24425 mRNA sequence 97.00 445908 R13580 Hs.13436 Hs.14623 Interferon, gamma-inducible protein 30 0.93 1.69 446291 BE397753 5 Hs.156672 1.00 106.00 446917 AI347863 447261 NM_006691 Hs.17917 extracellular link domain-containing 1 0.40 47.20 1.00 0.05 100.00 447432 AW958473 Hs.301957 nudix (nucleoside diphosphate linked mol 8.21 KIAA1233 protein 447482 AB033059 Hs.18705 Hs.29792 0.02 5.42 447997 ESTs, Weakly similar to I38022 hypotheti H00656 10 Hs.20887 hypothetical protein FLJ10392 1.00 79.00 448299 AA497044 0.42 448782 AL050295 Hs.22039 KIAA0758 protein 1.56 11.33 450575 NM_005859 Hs.29117 purine-rich element binding protein A AA040403 1.00 450584 Hs.60371 ESTs AW450461 Hs.203965 1.00 91.00 450693 **ESTs** 15 Al266484 450715 Hs.31570 ESTs, Weakly similar to KIAA1324 protein 1.00 152.00 1.00 86.00 DKFZP564D206 protein novel SH2-containing protein 3 Hs.25956 Hs.26054 451103 R52804 1.30 0.60 AF124251 451220 0.54 1.91 Z43948 Hs.326444 cartilage acidic protein 1 451668 1.00 AW023595 Hs.232048 **ESTs** 67.00 452197 20 purine-rich element binding protein A 4.53 11.07 452331 AA598509 Hs.29117 0.72 452353 C18825 Hs.29191 epithelial membrane protein 2 453049 BE537217 Hs.30343 **FSTs** 1.00 68.00 Hs.279746 vanilioid receptor-like protein 1 0.83 1.70 453107 NM_016113 Homo sapiens cDNA FLJ11422 fis, clone HE 132.00 453355 AW295374 Hs.31412 25 1.00 72.00 453390 453531 AA862496 Hs.28482 **ESTs** ESTs, Weakly similar to JC5795 CDEP prot 1.00 68.00 AA417940 gb:CM2-HT0342-091299-050-b05 HT0342 Homo up-regulated by BCG-CWS 2.89 454741 BE154396 0.57 82.00 Hs.284205 1.00 456579 AA287827 Hs.114727 1.98 Homo sapiens, clone MGC:16327, mRNA, com 0.79 456672 AK002016 30 Hs.252549 1.03 3.25 AF032906 cathepsin Z 457400 ESTs, Weakly similar to ALU4_HUMAN ALU S 1.00 113.00 457718 F18572 Hs.22978 544.00 qb:HSC1KA072 normalized infant brain cDN 1.00 459696 F03027 TABLE 10B 35 Unique Eos probeset identifier number CAT number: Gene cluster number Genbank accession numbers Accession: 40 Pkey 408074 CAT Number R20723 AA263003 AA333976 AA334725 AA334151 AW965490 AA310513 AI810530 D31302 AW134897 AA830127 AA046953 AI668930 103684_1 BE160198 AW935898 T11520 AW935930 AW856073 AW861034 411667 1253334_1 BE146973 BE146972 BE147042 BE147018 BE146783 BE147020 BE146781 BE147019 BE146766 BE147021 BE146952 BE146767 BE147044 413533 1375344_1 45 BE146797 BE146776 BE146985 BE146793 BE146768 BE146771 BE146954 BE146760 BE147048 BE147025 BE147030 AJ012074 U11087 L13288 X75299 L20295 AW630780 H14880 T28037 AJ872991 R72136 AW449839 T81622 T75697 T29519 R94105 T83923 R73300 AJ797007 R73390 AA981010 H74168 AJ689932 BE045543 AJ808418 AJ608912 AJ806573 AW884084 AW872978 AW872985 AA565655 AJ022915 R50647 R73210 H45098 R46451 AW166269 T71132 AJ264547 R52146 AJ304920 R73391 AW884059 AW884085 H73241 T60038 423387 22779_1 T79612 R73145 R50549 A1094557 A1668793 R72302 A1564366 W01956 AA418962 W32571 R72840 H45409 R72085 R46356 R46758 50 AA508805 AA418798 T83751 R94072 T16182 AA928785 AA903896 Z92546 AA330586 AI570568 AW341487 AI827050 AW298668 AI792189 AI015693 AI733599 AI572251 AI672488 AW193262 AI244716 AI864375 AI206100 AA912444 AI269365 AI640254 AW772466 AI867336 AA627604 H16914 AA358477 AA338009 423696 23112_1 430212 314437_1 AA469153 AI718503 AA469225 436532 421802_1 AA721522 AW975443 T93070 55 453531 97026_1 AA417940 AA036735 T07025 BE154396 AW817959 BE154393 454741 1232559 1 TABLE 10C 60 Unique number corresponding to an Eos probeset Pkev: Sequence source. The 7 digit numbers in this column are Genbank Identifier (GI) numbers. "Dunham I. et al." refers to the publication entitled "The DNA sequence of human chromosome 22." Dunham I. et al., Nature (1999) 402:489-495. Indicates DNA strand from which exons were predicted. Strand: 65 Indicates nucleotide positions of predicted exons. Nt position: Pkey 400754 Strand Nt position 144559-144684 7331445 Plus 401045 8117619 Plus 90044-90184,91111-91345 70 401083 3242744 Plus 33192-33360 53526-53628,55755-55920,57530-57757 402474 7547175 Minus 114964-115136,115461-115585,115931-116047,117666-117771,118004-118102 6456148 Minus 402808 403021 7547270 120799-120966 126609-126773,139986-140205 403421 403438 9665041 Minus 75 90792-90938 Plus 9719679 9009-9534 403687 7387384 Plus 118692-118853 7717105 Minus 403764 91665-91946 404277 1834458 Minus 2769644 Plus 3512-3691 40428B 80 404394 3135305 Minus 37121-37205,37491-37762,41053-41140,41322-41593,41773-41919 404518 8151988 Pius 84494-84603 91057-91188 404916 7341826 Plus 80877-81418 8079395 Minus 405106

73121-73273

7636-8054

405257

405381

85

7329310

6006920

Minus

Table 11A shows about 84 genes upregulated in lung adenocarcinomas relative to other lung tumors, non-malignant lung disease, and normal lung. These genes were selected from about 59680 probesets on the Eos/Affyrnetrix Hu03 Genechlp array.

Table 11B show the accession numbers for those Pkey's lacking UnigenelD's for table 11A. For each probeset we have listed the gene cluster number from which the oligonucleotides were designed. Gene clusters were compiled using sequences derived from Genbank ESTs and mRNAs. These sequences were clustered based on sequence similarity using Clustering and Alignment Tools (DoubleTwist, Oakland California). The Genbank accession numbers for sequences comprising each cluster are listed in the "Accession" column.

Table 11C show the genomic positioning for those Pkey's lacking Unigene ID's and accession numbers in lable 11A. For each predicted exon, we have listed the genomic sequence source used for prediction. Nucleotide locations of each predicted exon are also listed.

Unique Eos probeset identifier number 15 Exemplar Accession number, Genbank accession number

UnigenelD: Unigene number Unigene gene title

5

10

Unigene Title: R1: Average of lung tumors (including squamous cell carcinomas, adenocarcinomas, small cell carcinomas, granulomatous and carcinoid tumors) divided by the

average of normal lung samples

Average of non-malignant lung disease samples (including bronchitis, emphysema, fibrosis, atelectasis, asthma) divided by the average of normal lung samples 20 R2:

		_		•		
	Pkey	ExAccn	UnigeneiD	Unigene Title	R1	R2
	403329			Target Exon	1.00	61.00
25	406399			NM_003122*:Homo saplens serine protease	1.00	39.00
25	406690	M29540	Hs.220529	carcinoembryonic antigen-related cell ad	226.37 0.77	350.00 1.18
	407869	A1827976	Hs.24391	hypothetical protein FLJ13612	1.00	10.00
	407881 408908	AW072003 BE296227	Hs.40968 Hs.250822	heparan sulfate (glucosamine) 3-O-sulfot serine/threonine kinase 15	7.76	1.00
•	409103	AF251237	Hs.112208	XAGE-1 protein	80.44	40.00
30	409187	AF154830	Hs.50966	carbamoyl-phosphale synthetase 1, mitoch	1.00	1.00
	409269	AA576953	Hs.22972	hypothetical protein FLJ13352	1.00	1.00
	410076	T05387	Hs.7991	ESTs	1.12	1.50
	410102	AW248508	Hs.279727	Homo sapiens cDNA FLJ14035 fis, clone HE	9.89	1.00
	410399	BE068889		synuclein, gamma (breast cancer-specific	0.92	1.06
35	411908	L27943	Hs.72924	cytidine deaminase	1.00	1.00
	412612	NM_000047	Hs.74131	arylsulfatase E (chondrodysplasia puncta	1.02	1.03
	414075	U11862	Hs.75741	amiloride binding protein 1 (amine oxida	0.84	1.07
	416208	AW291168	Hs.41295	ESTs, Weakly similar to MUC2_HUMAN MUCIN	3.67	1.00
40	417542	J04129	Hs.82269	progestagen-associated endometrial prote	1.28	1.35 1.00
40	419183	U60669	Hs.89663	cytochrome P450, subfamily XXIV (vitamin	1.00 13.05	115.00
	419502	AU076704	Hs.303154	fibrinogen, A alpha polypeptide popeye protein 3	1.00	13.00
	419631 420931	AW188117 AF044197	Hs.100431	small inducible cytokine B subfamily (Cy	1.00	8.00
	421155	H87879	Hs.102267	lysyl oxidase	1.00	15.00
45	421190	U95031	Hs.102482	mucin 5, subtype B, tracheobronchial	1.17	1.55
10	421474	U76362	Hs.104637	solute carrier family 1 (glutamate trans	1.46	1.76
	421515	Y11339	Hs.105352	GalNAc alpha-2, 6-sialytransferase 1, 1	1.00	3.00
	421582	Al910275		trefoll factor 1 (breast cancer, estroge	1.23	1.00
	422026	U80736	Hs.110826	trinucleotide repeat containing 9	1.00	52.00
50	422095	AI868872	Hs.282804	hypothetical protein FLJ22704	4.37	2.34
	422311	AF073515	Hs.114948	cytokine receptor-like factor 1	1.15	1.78
	422867	L32137	Hs.1584	cartilage oligomeric matrix protein (pse	1.69	3.17
	423472	AF041260	Hs.129057	breast carcinoma amplified sequence 1	48.13	72.00
55	423554	M90516	Hs.1674	glutamine-fructose-6-phosphate transamin	1.00	50.00
55	424502	AF242388	Hs.149585	lengsin	1.00 1.00	1.00 59.00
	424544	M88700	Hs.150403	dopa decarboxylase (aromatic L-amino aci NIMA (never in mitosis gene a)-related k	21.35	1.00
	424905 424960	NM_002497 BE245380	Hs.153704 Hs.153952	5' nucleotidase (CD73)	1.00	1.00
	425523	AB007948	Hs.158244	KIAA0479 protein	1.00	35.00
60	426230	AA367019	Hs.241395	protease, serine, 1 (trypsin 1)	1.00	83.00
•	427701	AA411101	Hs.243886	nuclear autoantigenic sperm protein (his	7.41	34.00
	428585	AB007863	Hs.185140	KIAA0403 protein	1.00	6.00
	428758	AA433988	Hs.98502	hypothetical protein FLJ14303	1.06	1.13
	429170	NM_001394	Hs.2359	dual specificity phosphatase 4	16.18	105.00
65	429263	AA019004	Hs.198396	ATP-binding cassette, sub-family A (ABC1	1.07	1.00
	429610	AB024937	Hs.211092	LUNX protein; PLUNC (palate lung and nas	1.59	1.69
	430508	AI015435	Hs.104637	ESTs	4.75	7.27
	430985	AA490232	Hs.27323	ESTs, Weakly similar to 178885 serine/th	0.94 5.66	1.28 15.00
70	431548	AI834273	Hs.9711	novel protein J domain containing protein 1	49.76	37.00
70	431566	AF176012 AA536130	Hs.260720 Hs.149018	Novel human gene mapping to chomosome 20	1.19	1.47
	431986 432375	BE536069	Hs.2962	S100 calcium-binding protein P	1.65	1.06
	432677	NM_004482	Hs.278611	UDP-N-acetyl-aipha-D-galactosamine:polyp	1.00	48.00
	433556	W56321	Hs.111460	calcium/calmodulin-dependent protein kin	1.00	19.00
75	433819	AW511097	Hs.112765	ESTs	3.71	8.00
• -	434001	AW950905	Hs.3697	serine (or cysteine) proteinase inhibito	29.31	72.00
	434424	Al811202	Hs.325335	Homo sapiens cDNA: FLJ23523 fis, clone L	1.00	64.00
	434792	AA649253	Hs.132458	ESTs	8.52	44.00
00	436217	T53925	Hs.107	fibrinogen-like 1	57.97	31.00
80	436749	AA584890	Hs.5302	lectin, galactoside-binding, soluble, 4	1.10	1.41
	436972	AA284679	Hs.25640	claudin 3	1.59	1.46
	437866	AA156781	II. F040	metallothionein 1E (functional)	3.62	101.00
	437935	AW939591	Hs.5940	mucin 13, epithelial transmembrane Williams-Beuren syndrome chromosome regi	1.60 1.00	1.39 1.00
85	438915 439451	AA280174	Hs.285681 Hs.278554	heterochromatin-like protein 1	23.28	52.00
33	403451	AF086270	∩3.£/0004	Heres continue trace biorent i	20.20	02.00

	w	O 02/086	443				PCT/US02/12476
	439759 441031 441377 443614	AL359055 Al110684 BE218239 AV655386	Hs.67709 Hs.7645 Hs.202656 Hs.7645	Homo saplens mRNA full length insert cDN fibrinogen, B beta polypepiide ESTs fibrinogen, B beta polypepiide	1.00 1.41 22.03 1.00	21.00 99.00 1.00 16.00	
5	443813 443991 444670 444931	AA876372 NM_002250 H58373 AV652066	Hs.93961 Hs.10082 Hs.332938 Hs.75113	Homo sapiens mRNA; cDNA DKFZp667D095 (fr potassium intermediate/smail conductance hypothetical protein MGC5370 general transcription factor IIIA	1.20 5.71 1.98 1.00	1.99 6.87 38.00 54.00	
10	446102 446163 446469 447388	AW168067 AA026880 BE094848 AW630534	Hs.317694 Hs.25252 Hs.15113 Hs.76277	ESTs Homo sapiens cDNA FLJ13603 fis, clone PL homogentisate 1,2-dioxygenase (homogenti Homo sapiens, clone MGC:9381, mRNA, comp	1.00 1.00 1.00 1.24	1.00 36.00 11.00 1.16	
15	447532 448243 448844 449444 451807	AK000614 AW369771 AI581519 AW818436 W52854	Hs.18791 Hs.52620 Hs.177164 Hs.23590	hypothetical protein FLJ20607 integrin, beta 8 ESOlute carrier family 16 (monocarboxylic hypothetical protein FLJ23293 similar to	1.23 15.84 1.00 1.00 1.55	1.63 1.00 31.00 83.00 35.00	
20	452689 453392 453464 453735	F33868 U23762 Al884911 Al066629	Hs.284176 Hs.32964 Hs.32989 Hs.125073	transferrin SRY (sex determining region Y)-box 11 receptor (calcitonin) activity modifying ESTs	1.54 - 1.00 1.55 1.01	1.44 16.00 2.45 1.30	
	TABLE 11	18					
25	Pkey: CAT num Accession	ber: Gene clust	s probesetide ter number accession num				
30	Pkey 410399	CAT Numbe 11995_1	BE068 Al9365	889 BE068882 AF044311 AF017256 NM_003087 A 27 AA804675 AA394097 Al139933 AA946606 BE1	71313 AA722407	7 AA293803 A14	AA872836 BE298825 BE299889 AID16464 AI684600 68480 AA056035 AA055968 AW796957 AI637713 421708 AW265211 AI493266 AA366132 AW966044
35	419502	18535_1	AU076 T68367 T68220 T73203 T74756	704 T74854 T74860 T72098 T73265 T73873 T6916 7 T68401 T53959 T72360 T72099 T60377 T58961 T 0 T74673 T71800 T68355 T61227 T62738 T69317 T	0	36 T60385 T734 164738 T74645 173768 T73962 445 T60424 T6 168607 T58898	10 T68781 T67845 T67593 T73952 T67864 T60630 T72037 T68688 T72063 T73258 T72826 T64242 T73382 T68914 T70975 T73400 T60631 T73277 7922 T67736 T68716 T67765 T74765 T73819 T58719 T64305 T71908 T68107
40			N3359- AA312 AW470 AA235	4 AA344542 AW805054 Al207457 T61743 AA0267: 91777 A40156 H66239 AV652989 H38728 R88521 AV 91774 AV651256 N54417 AA812862 AW182929 A111 252 T27853 T47778 R95746 H70620 AA701463 AW	17 H94389 AA38 1655200 R95790 1192 H61463 H 1827166 R98475	2695 AA918409 W03250 W009 72060 AA34450 C20925 AV657	13 AA344136 AV660126 R97923 AA343596 13 H38639 Al277511 AV661108 Al207625 T47810 1287 T71959 T71313 T73920 T73333 T61618 T69293
45			AA344 T70475 T72517 T69366	7 R02292 T60599 T69206 T70452 T74677 R29366 3 T69358 T68258 AV650429 T73341 T61702 T7459	4 AA343853 T73 18 A1110639 AA3 161277 T74914 18 T40095 K0227	3909 T68070 T7 344603 AF0635 T60352 R29675 72 T40106 AA34	'2065 H72149 T73493 T73495 AV645993 R02293 13 T64696 T68516 T72223 T60507 T67633 R29500 174843 AV645792 AA344408 T69197 T72057 13045 AA341908 AA341907 AA342807 AA341964
50			AA345 AA344 AA693	r 772042 762764 A1064899 AA343080 767832 772 234 767598 AA011414 768036 H48262 A1207557 7 583 760362 H58121 795711 772803 768055 77717 592 A1248502 R29454 764764 757001 773052 771 r 773317 774273 769420 768245 774380 767862 7	68219 W86031 5 R29036 T727 429 T51176 T58	T69081 T64232 93 T69122 T645	
55	421582	2041_1	Al9102 AA568 BE074	75 X00474 X52003 X05030 NM_003225 AA314326 312 AA614409 AA307578 AI925552 AW950155 AI9	AA308400 AA5 10083 M12075 I 009769 AW050	3E074052 AW04 690 AA858276 (R55389 A1001051 AW050700 AW750216 AA614539
	437866	44433_2	AA156		3977 AW44490	4 AW602574 BE	E164040 BE164012 BE163972 BE163974 BE163992
60	451807	8865_1	W5285	489 AW874142 A1471883 W84421 AA166860 4 AL117600 BE208116 BE208432 BE206239 BE08 662 AW449519 AA993634 A1806539 AA351618 AV			
65	TABLE 1	1C					
	Pkey: Ref:	Sequence	source. The 7	nding to an Eos probeset * digit numbers in this column are Genbank Identifier mosome 22.* Dunham I. et al., Nature (1999) 402:4	(GI) numbers. '	'Dunham I. et al	refers to the publication entitled "The DNA
70	Strand: Nt_position	Indicates C	NA strand from	m which exons were predicted. tions of predicted exons.			

Pkey 403329 406399

75

Ref 8516120 9256288 Strand Plus Minus Nt_position 98450-96598 63448-63554

TABLE 12A: Genes Distinguishing Squamous Cell Carcinoma from Other Lung Diseases and Normal Lung

Table 12A shows about 72 genes upregulated in squamous cell carcinomas of the lung relative to other lung tumors, non-malignant lung disease, and normal lung. These genes 5 were selected from about 59680 probesets on the Eos/Affymetrix Hu03 Genechip array.

Table 12B show the accession numbers for those Pkey's lacking UnigenelD's for table 12A. For each probaset we have listed the gene cluster number from which the oligonucleotides were designed. Gene clusters were compiled using sequences derived from Genbank ESTs and mRNAs. These sequences were clustered based on sequence similarity using Clustering and Alignment Tools (DoubleTwist, Oakland California). The Genbank accession numbers for sequences comprising each cluster are listed in the

Table 12C show the genomic positioning for those Pkey's lacking Unigene ID's and accession numbers in table 12A. For each predicted exon, we have listed the genomic sequence source used for prediction. Nucleotide locations of each predicted exon are also listed.

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Pkey: ExAccn:

Unique Eos probeset identifier number Exemplar Accession number, Genbank accession number

UnigenelD: Unigene number

Unigens gene tille

Average of lung tumors (including squamous cell carcinomas, adenocarcinomas, small cell carcinomas, granulomatous and carcinoid tumors) divided by the Unigene Title: 20 R1:

average of normal lung samples

Average of non-malignant lung disease samples (Including bronchitis, emphysema, fibrosis, atelectasis, asthma) divided by the average of normal lung samples R2:

	R2:	Average	or non-maugn	ant lung disease samples (including bronchius, emphy	seilla, iiulusis, a	iciociasis, esi
	Pkey	ExAcon	UnigenelD	Unigene Tille	R1	R2
25	400289	X07820	Hs.2258	matrix metalloproteinase 10 (stromelysin	132.45	4.00
	400666			NM_002425:Homo saplens matrix metallopro	3.26	3.22
	401780			NM_005557*:Homo sapiens keratin 16 (foca	26.47	10.50
	401781			Target Exon	10.33	4.61
	401785			NM_002275*:Homo saplens keratin 15 (KRT1	4.13	2.70
30	401994			Target Exon	61.84	47.00
	402075			ENSP00000251056*:Plasma membrane calcium	1.00	1.00
	404996			Target Exon	1.00 173.91	1.00 108.00
	407839	AA045144	Hs.161566	ESTs	151.17	8.00
35	408000	L11690	Hs.620 Hs.46320	bullous pemphigoid antigen 1 (230/240kD) Small proline-rich protein SPRK [human,	1.98	1.24
22	408522 410561	AI541214 BE540255	Hs.6994	Homo saplens cDNA: FLJ22044 fis, clone H	10.04	1.00
	415091	AL044872	Hs.77910	3-hydroxy-3-methylgiutaryl-Coenzyme A sy	1.00	30.00
	415817	U88967	Hs.78867	protein tyrosine phosphatase, receptor-t	24.30	1.00
	416658	U03272	Hs.79432	fibrillin 2 (congenital contractural ara	53.29	51.00
40	417034	NM_006183	Hs.80962	neurotensin	1.00	1.00
••	417366	BE185289	Hs.1076	small proline-rich protein 1B (cornifin)	8.97	3.27
	418663	AK001100	Hs.41690	desmocollin 3	112.17	19.00
	418678	NM_001327	Hs.87225	cancer/testis antigen	1.18	1.10 .
	419121	AA374372	Hs.89626	parathyroid hormone-like hormone	1.00	1.00
45	420783	AI659838	Hs.99923	lectin, galactoside-binding, soluble, 7	3.04	1.25
	421773	W69233	Hs.112457	ESTs	1,12	1.14
	421948	L42583	Hs.334309	keratin 6A	51.83	20.25
	421978	AJ243662	Hs.110196	NICE-1 protein	1.01	0.91
50	422158	L10343	Hs.112341	protease inhibitor 3, skin-derived (SKAL	2.37	1.10
50	422440	NM_004812	Hs.116724	aldo-keto reductase family 1, member B10	47.53 76.00	32.00
	423634	AW959908	Hs.1690	heparin-binding growth factor binding pr	76.02	1.00 1.00
	423725	AJ403108	Hs.132127	hypothetical protein LOC57822	4.20 10.14	51.00
	423738	AB002134	Hs.132195	airway trypsin-like protease	233.42	68.00
55	424012	AW368377	Hs.137569 Hs.138202	tumor protein 63 kDa with strong homolog serine (or cysteine) proteinase inhibito	1.00	1.00
55	424046 424098	AF027866 AF077374	Hs.139322	small proline-rich protein 3	137.82	54.00
	424834	AK001432	Hs.153408	Homo sapiens cDNA FLJ10570 fis, clone NT	56.19	12.00
	425650	NM_001944	Hs.1925	desmoglein 3 (pemphigus vuigaris antigen	33.45	1.00
	427099	AB032953	Hs.173560	odd Oz/ten-m homolog 2 (Drosophlia, mous	4.24	17.00
60	427335	AA448542	Hs.251677	G antigen 7B	51.83	4.00
	428182	BE386042	Hs.293317	ESTs, Weakly similar to GGC1_HUMAN G ANT	1.00	1.00
	428645	AA431400	Hs.98729	ESTs, Weakly similar to 2017205A dihydro	1.00	16.00
	428748	AW593206	Hs.98785	Ksp37 protein	1.00	87.00
	429259	AA420450	Hs.292911	ESTs, Highly similar to S60712 band-6-pr	2.01	1.18
65	429538	BE182592	Hs.11261	small proline-rich protein 2A	4.43	2.90
	429903	AL134197	Hs.93597	cyclin-dependent kinase 5, regulatory su	11.80	1.00
	430486	BE062109	Hs.241551	chloride channel, calcium activated, fam	12.28	41.00
	430890	X54232	Hs.2699	glypican 1	1.58	1.40
70	431009	BE149762	Hs.48956	gap junction protein, beta 6 (connexin 3	60.25	28.00 2.51
70	431846	BE019924	Hs.271580	uroplatin 18	4.49 1.20	1.09
	433091	Y12642	Hs.3185	lymphocyte antigen 6 complex, locus D ESTs	40.98	27.00
	434360	AW015415	Hs.127780	cytochrome P450, subfamily IVF, polypept	1.00	1.00
	434880	U02388	Hs.101 Hs.211238	interleukin-1 homolog 1	1.00	38.00
75	435505 435793	AF200492 AB037734	Hs.4993	KIAA1313 protein	23.68	42.00
13	436511	AA721252	Hs.291502	ESTs	16.76	14.00
	438403	AA806607	Hs.292206	ESTs	1.00	1.00
	439285	AL133916		hypothetical protein FLJ20093	46.23	139.00
	439606	W79123	Hs.58561	G protein-coupled receptor 87	33.61	1.00
80	439670	AF088076	Hs.59507	ESTs, Weakly similar to AC004858 3 U1 sm	1.00	1.00
	439706	AW872527	Hs.59761	ESTs, Weakly similar to DAP1_HUMAN DEATH	86.55	11.00
	440325	NM_003812	Hs.7164	a disintegrin and metalloproteinase doma	62.88	147.00
	441525	AW241867	Hs.127728	ESTs	1.53	1.42
0.5	443162	T49951	Hs.9029	DKFZP434G032 protein	31.11	38.00
85	444378	R41339	Hs.12569	ESTs	1.00	1.00

	WC) 02/086 4	143				PCT/US02/12476	
5	446292 447078 447342 449003 449101 450832 452240 453317 453830 454098	AF081497 AW885727 AI199268 X76342 AA205847 AW970602 AI591147 NM_002277 AA534296 W27953 AI368680	Hs.279682 Hs.9914 Hs.19322 Hs.389 Hs.23016 Hs.105421 Hs.61232 Hs.41696 Hs.20953 Hs.292911 Hs.816	Rh type C glycoprotein ESTs Homo septens, Similar to RIKEN cDNA 2010 alcohol dehydrogenase 7 (class IV), mu o G protein-coupled receptor ESTs ESTs keratin, hair, acidic,1 ESTs ESTs, Highly similar to S60712 band-6-pr SRY (sex determining region Y)-box 2	1.55 47.24 28.63 1.00 2.58 25.17 13.42 1.19 24.92 1.26 206.11	1.26 24.00 1.00 1.00 27.00 36.00 1.00 1.27 25.00 1.11	·	
	TABLE 12B							
15	Pkey: CAT numbe Accession:	r. Gene cluste	probeset iden r number cession numb					
20 .		CAT Number 47085_1	Accession AL13391 AA7755	on 16 N79113 AF086101 N76721 AW950828 AA364013 52 N62351 N59253 AA626243 Al341407 BE175639	3 AW955684 Al34 AA456968 Al3589	6341 AI867454 N54784 918 AA457077	4 Al655270 Al421279 AW014882	
25	TABLE 12C .							
23	Pkey: Ref:							
30	Strand: Nt_position:	Indicates DI Indicates nu	NA strand from	which exons were predicted. ons of predicted exons.				
35	Pkey 400666 401780 401781 401785 401994 402075 404996	Ref 8118496 7249190 7249190 7249190 4153858 8117407 6007890	Strand Plus Minus Minus Minus Minus Minus Plus Plus	Nt_position 17982-18115,20297-20456 28397-28617,28920-29045,29135-29296,29411-2 83215-83435,83531-83656,83740-83901,84237-8 165776-165996,166189-166514,166408-166569, 42904-43124,43211-43336,44607-44763,45199-4 121907-122035,122804-122921,124019-124161,1 37999-38145,38652-38998,39727-39872,40557-4	4393,84955-8503 167112-167268,16 5281,46337-4673 124455-124610,12	7,86290-86814 37387-167469,168634- 2 25672-126076	168942	
40								

TABLE 13A: Genes Distinguishing Non-Malignant Lung Disease from Lung Tumors and Normal lung

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402075

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8117407

Table 13A shows about 23 genes upregulated in non-malignant lung disease relative to lung tumors and normal lung. These genes were selected from about 59680 probesets on the Eos/Affymetrix Hu03 Genechip array.

Table 13B show the accession numbers for those Pkey's lacking UnigenelD's for table 13A. For each probeset we have listed the gene cluster number from which the oligonucleotides were designed. Gene clusters were complied using sequences derived from Genbank ESTs and mRNAs. These sequences were clustered based on sequence similarity using Clustering and Alignment Tools (DoubleTwist, Oakland California). The Genbank accession numbers for sequences comprising each cluster are listed in the "Accession" column.

Table 13C show the genomic positioning for those Pkey's lacking Unigene ID's and accession numbers in table 13A. For each predicted exon, we have listed the genomic sequence source used for prediction. Nucleotide locations of each predicted exon are also listed.

						•				
15	Pkev:	Uniqu	e Fos probes	et identifier number						
	ExAccn:	Exem	olar Accession	n number, Genbank accession number						
	UnigenelD:		ne number	•		•				
	Unigene Title	e Unine	Lintrene dene title							
	R1:	Avera	ge of lung tun	nors (including squamous cell carcinomas, adenocar	(including squamous cell carcinomas, adenocarcinomas, small cell carcinomas, granulomatous and carcinoid tumors) divided by the					
20		avera	ge of normal l	ung samples	ng samples					
	R2:	Avera	ge of non-mal	nant lung disease samples (including bronchitls, emphysema, fibrosis, atelectasis, asthma) divided by the average of normal lung samples						
						•				
		xAccn	UnigenelD	Unigene Title	R1	R2.				
26		1436323	Hs.31141	Homo sapiens mRNA for KIAA1568 protein,	1.00	230.00				
25		A376836	Hs.76728	ESTs	1.00 1.00	128.00 173.00				
		65998	Hs.285243	hypothetical protein FLJ22029	1.00	145.00				
		20350 L049176	Hs.78913 Hs.82223	chemokine (C-X3-C) receptor 1 chordin-like	1.00	179.00				
		A228776	Hs.191721	ESTs	1.00	140.00				
30		20893	Hs.325823	ESTs, Moderately similar to ALU5_HUMAN A	1.00	156.00				
50	422000 K	A464840	Hs.131987	ESTs	1.00	167.00				
		89832	Hs.170278	ESTS	1.00	141.00				
		A453800	Hs.192793	ESTS	1,00	138.00				
		A488988	Hs.293796	ESTs	1,00	133.00				
35		E041395	113.200100	ESTs, Weakly similar to unknown protein	23.32	941.00				
55	431385 B	E178536	Hs.11090	membrane-spanning 4-domains, subfamily A	1.00	157.00				
			Hs.268107	multimerin	1.00	157.00				
		A721522	***************************************	gb:nv54h12.r1 NCI_CGAP_Ew1 Homo sapiens	1.00	218.00				
		1669586	Hs.222194	ESTs	1.00	147.00				
40	438202 A	W169287	Hs.22588	ESTs	1.00	141.00				
			Hs.101689	ESTs	1.00	167.00				
		L120214	Hs.7117	glutamate receptor, ionotropic, AMPA 1	1.00	151.00				
		25899	Hs.201591	ESTs	1.00	141.00				
40		67837	Hs.169872	ESTs	1.00	116.00				
45		1000341	Hs.220491	ESTs	1.00	192.00				
	459587 A	A031956		gb:zk15e04.s1 Soares_pregnant_uterus_NbH	1.00	154.00				
	T101 F 400									
	TABLE 13B									
50	Pkev:	Deleve E	on nombocat in	ientifier number						
50	CAT number			retibliet itatioet		•				
	Accession:		accession nu	mhers .						
		Cinbank	600000011111							
	Pkey	CAT Num	ber Accessio	on.						
55	431089	327825_1		95 AA491826 AA621946 AA715980 AA666102						
	436532	421802_1		22 AW975443 T93070						
		_								
	TABLE 13C									
60										
	Pkey:	Unique ni	umber correst	ponding to an Eos probeset		A A A A A A A A A A A A A A A A A A A				
	Ref:	Sequence	source. The	7 digit numbers in this column are Genbank Identifi	er (GI) numbers. "D	unham I, et al." refers to the publication entitled "The DNA				
				romosome 22." Dunham I. et al., Nature (1999) 402	489-495.					
65	Strand:			rom which exons were predicted.						
65	Nt_position:	indicates	nucleotide po	sitions of predicted exons.						
	Die	Def	Circus	Alt modilion						
	Pkey	Ref	Strand	Nt_position						

121907-122035,122804-122921,124019-124161,124455-124610,125672-126076

Table 14A shows the subcellular localization and preferred utility for the genes appearing in Tables 9A and 10A. mAb symbolizes monoclonal antibody, diag symbolizes diagnostic, s.m. symbolizes small molecule, and CTL symbolizes cytotoxic lymphocytic ligand. These genes were selected from 59680 probesets on the Eos/Affymetrix Hu03 Genechip array. 5

Table 14B show the accession numbers for those Pkey's lacking UnigeneiD's for table 14A. For each probeset we have listed the gene cluster number from which the oligonucleotides were designed. Gene clusters were compiled using sequences derived from Genbank ESTs and mRNAs. These sequences were clustered based on sequence similarity using Clustering and Alignment Tools (DoubleTwist, Oakland California). The Genbank accession numbers for sequences comprising each cluster are listed in the "Accession" column.

Table 14C show the genomic positioning for those Pkey's lacking Unigene ID's and accession numbers in table 14A. For each predicted exon, we have listed the genomic sequence source used for prediction. Nucleotide locations of each predicted exon are also listed.

Pkey: Unique Eos probeset identifier number
ExAcon: Exemplar Accession number, Genbank accession number
UnigenelD: Unigene number

Unigene Title: Unigene gene title
Pref.Utility: Preferred Utility
Pred.Loc: Preferred utility

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	, , , ,					
	Pkey	ExAccn	UnigenelD	Unigene Title	Pref Utility	Pred. Loc
	400289	X07820	Hs.2258	matrix metalloproteinase 10 (stromelysin	mAb & diag & s.m.	extracellular
25	400303	AA242758	Hs.79136	LIV-1 protein, estrogen regulated	mAb	plasma membrane
	402075		,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	ENSP00000251056*:Plasma membrane calcium	mAb & diag	secreted
	407811	AW190902	Hs.40098	cysteine knot superfamily 1, BMP antagon	diag	secreted
	408243	Y00787	Hs.624	interleukin 8	diag	secreted
	408790	AW580227	Hs.47860	neurotrophic tyrosine kinase, receptor,	mAb & s.m.	plasma membrane
30	408908	BE296227	Hs.250822	serine/threonine kinase 15	s.m.	cytoplasm
	409041	AB033025	Hs.50081	Hypothetical protein, XP_051860 (KIAA119	CTL & diag	secreted
	409103	AF251237	Hs.112208	XAGE-1 protein	CTL	nuclear
	409420	Z15008	Hs.54451	taminin, gamma 2 (nicein (100kD), kalini	diag	secreted
	409632	W74001	Hs.55279	serine (or cysteine) proteinase inhibito	diag	secreted
35	409757	NM_001898	Hs.123114	cystatin SN	diag	extracellular
	409893	AW247090	Hs.57101	minichromosome maintenance deficient (S.	CTL	nuclear
	409956	AW103364	Hs.727	inhibin, beta A (activin A, activin AB a	diag	extracellular
	410001	AB041036	Hs.57771	kallikrein 11	dlag	extracellular
	410407	X66839	Hs.63287	carbonic anhydrase IX	mAb & s.m.	plasma membrane
40	410418	D31382	Hs.63325	transmembrane protease, serine 4	mAb & diag & s.m.	plasma membrane
	412140	AA219691	Hs.73625	RAB6 interacting, kinesin-like (rabkines	s.m.	
	412719	AW018610	Hs.816	ESTs	s.m.	nuclear
	414774	X02419	Hs.77274	plasminogen activator, urokinase	diag	extracellular
	414883	AA926960		CDC28 protein kinase 1	s.m.	
45	415138	C18356	Hs.295944	tissue factor pathway inhibitor 2	CTL & dlag	extracellular
	415669	NM_005025	Hs.78589	serine (or cysteine) proteinase inhibito	mAb & diag & s.m.	secreted
	415817	U88967	Hs.78867	protein tyrosine phosphatase, receptor-t	mAb & s.m.	plasma membrane
	416658	U03272	Hs.79432	fibrillin 2 (congenital contractural ara	diag	extracellular
5 0	417034	NM_006183	Hs.80962	neurotensin	diag	extracellular
50	417079	U65590	Hs.81134	Interleukin 1 receptor antagonist	diag	extracellular
	417308	H60720	Hs.81892	KIAA0101 gene product	s.m.	mitochondrial
	417389	BE260964	Hs.82045	midkine (neurite growth-promoting factor	mAb & diag	secreted
	417433	BE270266	Hs.82128	5T4 oncofetal trophoblast glycoprotein	mAb	plasma membrana
55	417933	X02308	Hs.82962	thymidylate synthetase	s.m.	endoplasmic reticulum
55	418478	U38945	Hs.1174	cyclin-dependent kinase inhibitor 2A (me	s.m.	cytoplasm
	418506	AA084248	Hs.85339	G protein-coupled receptor 39	mAb & s.m.	plasma membrane
	418678	NM_001327	Hs.167379	cancer/testis antigen (NY-ESO-1)	CTL	cytoplasmic secreted
	419121	AA374372	Hs.89626	parathyroid hormone-like hormone	diag mAb & s.m.	plasma membrane
60	419171	NM_002846	Hs.89655	protein tyrosine phosphatase, receptor t	CTL & s.m.	mitochondrial
UU	419183 419216	U60669 AU076718	Hs.89663 Hs.164021	cytochrome P450, subfamily XXIV (vitamin small inducible cytokine subfamily B (Cy	diag	secreted
	419235	AW470411	Hs.288433	neurotrimin	mAb & diag	plasma membrane
	419452	U33635	Hs.90572	PTK7 protein tyrosine kinase 7	mAb & s.m.	plasma membrane
	419556	U29615	Hs.91093	chitinase 1 (chitotriosidase)	mAb & diag	extracellular*
65	420610	Al683183	Hs.99348	distal-less homeo box 5	CTL	nuclear
05	421110	AJ250717	Hs.1355	cathepsin E	sm & diag	extracellular
	421379	Y15221	Hs.103982	small inducible cytokine subfamily B (Cy	diag	secreted
	421474	U76362	Hs.104637	solute carrier family 1 (glutamate trans	mAb & s.m.	plasma membrane
	421552	AF026692	Hs.105700	secreted frizzled-related protein 4	diag	secreted
70	421753	BE314828	Hs.107911	ATP-binding cassette, sub-family B (MDR/	mAb & s.m.	plasma membrane
. •	421817	AF146074	Hs.108660	ATP-binding cassette, sub-family C (CFTR	mAb & s.m.	plasma membrane
	422109	S73265	Hs.1473	gastrin-releasing peptide	diag	secreted
	422158	L10343	Hs.112341	protease inhibitor 3, skin-derived (SKAL	diag	secreted
	422282	AF019225	Hs.114309	apolipoprotein L	diag	secreted
<i>75</i>	422283	AW411307	Hs.114311	CDC45 (cell division cycle 45, S.cerevis	s.m.	nuclear
	422424	Al186431	Hs.296638	prostate differentiation factor	diag	extracellular
	422765	AW409701	Hs.1578	baculoviral IAP repeat-containing 5 (sur	s.m.	cytoplasm
	422809	AK001379	Hs.121028	hypothetical protein FLJ10549	s.m.	nuclear
	422867	L32137	Hs.1584	cartilage oligomeric matrix protein (pse	diag	extracellular
80	422956	BE545072	Hs.122579	ECT2 protein (Epithelial cell transformi	CTL & s.m.	
-	423634	AW959908	Hs.1690	heparin-binding growth factor binding pr	diag	
	423673	BE003054	Hs.1695	matrix metalloproteinase 12 (macrophage	mAb & diag & s.m.	secreted
	423961	D13666	Hs.136348	periostin (OSF-2os)	mAb & diag	extracellular
0.5	424046	AF027866	Hs.138202	serine (or cysteine) proteinase inhibito	diag	secreted
85	424381	AA285249	Hs.146329	protein kinase Chk2	s.m.	nuclear

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	424502	AF242388	Hs.149585	lengsin	e m	cytopiasmic
				Integrin, alpha 5 (fibronectin receptor,	s.m. mAb & s.m.	plasma membrane
	424503 424687	NM_002205 J05070	Hs.149609 Hs.151738	matrix metalloproteinase 9 (gelatinase B	diag	extracellular
	425247	NM_005940	Hs.155324	matrix metalloproteinase 11 (stromelysin	mAb & diag & s.m.	secreted
5	425322	U63630	Hs.155637	protein kinase, DNA-activated, catalytic	s.m.	cytoplasmic
,	425650	NM_001944	Hs.1925	desmoglein 3 (pemphigus vulgaris antigen	mAb	plasma membrane
	425734	AF056209	Hs.159396	peptidylglycine alpha-amidating monooxyg	s.m.	
	425776	U25128	Hs.159499	parathyroid hormone receptor 2	mAb & diag	plasma membrane
	425852	AK001504	Hs.159651	death receptor 6, TNF superfamily member	mAb & s.m.	plasma membrane
10	426215	AW963419	Hs.155223	stanniocalcin 2	mAb & diag	secreted
	426427	M86699	Hs.169840	TTK protein kinase	CTL & s.m.	nuclear
	426514	BE616633	Hs.170195	bone morphogenetic protein 7 (osteogenic	mAb & dlag	secreted
	427335	AA448542	Hs.251677	G antigen 7B	CTL	cytoplasmic
	427747	AW411425	Hs.180655	serine/threonine kinase 12	s.m.	cytoplasmic
15	428242	H55709	Hs.2250	leukemia inhibitory factor (cholinergic -	diag	
	428330	L22524	Hs.2256	matrix metalloproteinase 7 (matrilysin,	mAb & diag & s.m.	extracellular
	428450	NM_014791	Hs.184339	KIAA0175 gene product	s.m.	nuclear
	428479	Y00272	Hs.334562	cell division cycle 2, G1 to S and G2 to	s.m.	nuclear
20	428484	AF104032	Hs.184601	solute carrier family 7 (cationic amino	mAb & s.m.	plasma membrane
20	428664	AK001666	Hs.189095	similar to SALL1 (sal (Drosophila)-like	CTL & s.m.	nuclear
	428698	AA852773	Hs.334838	KIAA1866 protein	mAb	extracellular
	428748	AW593206	Hs.98785	Ksp37 protein	diag	mitochodria*
	428758	AA433988	Hs.98502	CA125 antigen; mucin 16	diag diag	extracellular
25	428969	AF120274	Hs.194689	artemin gap junction protein, beta 5 (connexin 3	mAb & s.m.	plasma membrane
23	429211	AF052693	Hs.198249		mAb & s.m.	plasma membrane
	429263	AA019004	Hs.198396	ATP-blnding cassette, sub-family A (ABC1	diag	secreted
	429547 429610	AW009166 AB024937	Hs.99376 Hs.211092	ESTs LUNX protein; PLUNC (palate lung and nas	mAb & diag	secreted
	429903	AL134197	Hs.93597	cyclin-dependent kinase 5, regulatory su	s.m.	000000
30	430488	BE062109	Hs.241551	chloride channel, calcium activated, fam	mAb & s.m.	plasma membrane
50	431462	AW583672	Hs.256311	granin-like neuroendocrine peptide precu	diag	extracellular
	431515	NM_012152	Hs.258583	endothelial differentiation, lysophospha	mAb & s.m.	plasma membrane
	431846	BE019924	Hs.271580	uroplakin 1B	mAb & dlag	plasma membrane
	431958	X63629	Hs.2877	cadherin 3, type 1, P-cadherin (placenta	mAb & diag	plasma membrane
35	432201	AI538613 ·	Hs.298241	Transmembrane protease, serine 3	mAb & diag & s.m.	plasma membrane
	433001	AF217513	Hs.279905	clone HQ0310 PRO0310p1	s.m.	nuclear
	435505	AF200492	Hs.211238	interleukin-1 homolog 1	diag	secreted
	436481	AA379597	Hs.5199	HSPC150 protein similar to ubiquitin-con	s.m.	
4.0	437016	AU076916	Hs.5398	guanine monphosphate synthetase	s.m.	cytoplasm
40	437044	AL035864	Hs.69517	differentially expressed in Fanconi's an	CTL	er _.
	437789	AI581344	Hs.127812	ESTs, Weakly similar to T17330 hypotheti	CTL	nuclear
	437852	BE001836	Hs.256897	ESTs, Weakly similar to dJ365O12.1 [H.sa	mAb & s.m.	plasma membrane
	439223	AW238299	Hs.250618	UL16 binding protein 2	mAb a	plasma membrane
45	439477	W69813	Hs.58042	ESTs, Moderately similar to GFR3_HUMAN G	mAb & s.m. mAb & s.m.	plasma membrane
45	439606	W79123	Hs.58561	G protein-coupled receptor 87	mAb & s.m.	plasma membrane
	439738 440006	BE246502	Hs.9598	sema domain, immunoglobulin domain (lg), NALP2 protein; PYRIN-Containing APAF1-li	s.m.	nuclear
	441362	AK000517 BE614410	Hs.6844 Hs.23044	RAD51 (S. cerevisiae) homolog (E coli Re	s.m.	1100001
	442117	AW664964	Hs.128899	ESTs; hypothetical protein for IMAGE:447	mAb & s.m.	plasma membrane
50	443247	BE614387	Hs.333893	c-Myc target JPO1	CTL	extracellular*
-	443426	AF098158	Hs.9329	chromosome 20 open reading frame 1	CTL	
	443859	NM_013409	Hs.9914	follistatin	diag	extracellular
	444006	BE395085	Hs.10086	type I transmembrane protein Fn14	mAb	plasma membrane
	444371	BE540274	Hs.239	forkhead box M1	s.m.	nuclear
55	444381	BE387335	Hs.283713	ESTs, Weakly similar to S64054 hypotheti	diag	secreted
	444781	NM_014400	Hs.11950	GPI-anchored metastasis-associated prote	mAb & diag	plasma membrane
	445537	AJ245671	Hs.12844	EGF-like-domain, multiple 6	mAb & diag	secreted
	446619	AU076643	Hs.313	secreted phosphoprotein 1 (osteopontin,	diag	secreted
~	446921	AB012113	Hs.16530	small Inducible cytokine subfamily A (Cy	diag	extracellular
60	447033	Al357412	Hs.157601	EST8	CTL & diag	secreted
	447342	Al199268	Hs.19322	Homo sapiens, Similar to RIKEN cDNA 2010	CTL	alasma mambana
	448243	AW369771	Hs.52620	Integrin, beta 8	mAb & s.m	plasma membrane
	448844	A)581519	Hs.177164	ESTS	mAb & s.m. mAb	plasma membrane
65	449048	Z45051	Hs.22920	similar to S68401 (cattle) glucose induc	s.m.	cytoplasm
UJ	449722	BE280074 NM_001044	Hs.23960 Hs.406	cyclin B1 solute carrier family 6 (neurotransmitte	mAb & s.m.	plasma membrane
	450001 450375	AA009647	115.400	a disintegrin and metalloproteinase doma	mAb & diag & s.m.	plasma membrane
	450375 450701	H39960	Hs.288467	hypothetical protein XP_098151 (leucine-	mAb & diag	plasma membrana
	450983	AA305384	Hs.25740	ERO1 (S. cerevislae)-like	diag	secreted
70	451668	Z43948	Hs.326444	cartilage acidic protein 1	mAb & diag	plasma membrane
	452281	T93500	Hs.28792	Homo sapiens cDNA FLJ11041 fis, clone PL	diag	
	452401	NM_007115	Hs.29352	tumor necrosis factor, alpha-induced pro	diag	extracellular
	452747	BE153855	Hs.61460	Ig superfamily receptor LNIR	mAĎ	plasma membrane
	452838	U65011	Hs.30743	preferentially expressed antigen in mela	CTL	nuclear
75	453968	AA847843	Hs.62711	High mobility group (nonhistone chromoso	CTL & s.m.	nuclear
	457489	A1693B15	Hs.127179	cryptic gene	diag	secreted
	TARICA	4D				
	TABLE 1	40				

TABLE 14B

80

Pkey: Unique Eos probeset identifier number CAT number: Gene cluster number Accession: Genbank accession numbers

Pkey CAT Number Accession

	WO 02/086443			PC17US02/12476					
5	414883	15024_1	AA08243 AA29275 AA87203 R75953	0 AA926959 W76521 W24270 W21526 AA037172 BE267636 H83186 AA469909 N86396 AA001348 BE535736 AA081745 BE566245 6 H72525 H77575 N49786 W80565 H78746 BE569085 W04339 R98127 T55938 BE279271 AW960304 T29812 AA476873 BE297387 3 AA177048 NM_001826 X54941 BE314366 AA908783 AI719075 BE270172 BE269819 AA889955 AI204630 W25243 A1935150 99 W72395 T99630 A1422691 H98460 N31428 BE255916 H03265 A1857576 AA776920 AA910844 AA459522 AA293140 AW514667 AW662396 AA662522 A1865147 A1423153 AW262230 AA584410 AA563187 AW024595 AW069734 A1828996 AA282997 AA876046 22 AA527373 AW972459 A1831360 AA621337 AA100926 AA772418 AA594628 A1033892 W95096 A1034317 AA398727 A1085031					
10	•		AA64328 A1139549 A1494230 A149421 AA95434	A459432 A1041437 AA932124 AA627684 AA935829 A1004827 A1423513 A1094597 H42079 R54703 A1630359 AA617681 AA978045 0 W44561 A1991988 A1537692 A1090262 AA740817 A1312104 A1911822 AA416871 A1185409 AA129784 AA701623 A1075239 10 AA633648 A1339996 A1336880 AA399239 A1078708 A1085351 A1362835 A1346618 A1146955 A1989380 A1348243 N92892 AA765850 10 A1278887 AA962596 A1492600 W80435 AA001979 R97424 A1129015 N24127 AA157451 AA235549 AA459292 AA037114 AA129785 11 AW059601 AW886710 R92790 N59755 A1361128 AW589407 H47725 H97534 H48076 H48450 T99631 AW300758 H03431 R76789 12 H77576 R96823 A1457100 N92845 N49682 H42038 BE220698 BE220715 H99552 AA701624 N74173 R54704 H79520 H72923					
15	450375	83327_1	W95095 AA00964	BE261919 AA769633 AA480310 AA507454 AA910586 Al203723 AW104725 W25611 W25071 T88980 H03513 T77589 R99156 R97470 AA702275 T77551 AA911952 H82956 N83673 AA283672 7 AA131254 AA374293 AW954405 H04410 AW606284 AA151166 BE157467 BE157601 H04384 W46291 AW663674 H04021 H01532 3 H03231 H59605 H01642 AA852876 AA113758 AA626915 AA746952 Al161014 AA099554 R69067					
20	TABLE 14C								
20	Pkey: Ref:	Sequence so	urce. The 7 o	ting to an Eos probeset ligit nurmbers In this column are Genbank Identifier (GI) numbers. "Dunharn I. et al." refers to the publication entitled "The DNA osome 22." Dunharn I. et al., Nature (1999) 402:489-495.					
25	Strand: Nt_position:	Indicates DN/	dicates ONA strand from which exons were predicted. dicates NA strand from which exons were predicted. dicates nucleolide positions of predicted exons.						
	Pkey	Ref	Strand	Nt_position					
30	402075	8117407	Plus	121907-122035,122804-122921,124019-124161,124455-124610,125672-126076					

TABLE 15A: Information for all sequences in Table 16

Table 15A shows the Seq ID No, Pkey, ExAcon, UnigeneID, and Unigene Title for all of the sequences in Table 16.

Table 15B show the accession numbers for those Pkey's lacking UnigenelD's for table 15A. For each probeset we have listed the gene cluster number from which the oligonucleotides were designed. Gene clusters were compiled using sequences derived from Genbank ESTs and mRNAs. These sequences were clustered based on sequence similarly using Clustering and Alignment Tools (DoubleTwist, Oakland California). The Genbank accession numbers for sequences comprising each cluster are listed in the 5

Table 15C show the genomic positioning for those Pkey's tacking Unigene ID's and accession numbers in table 15A. For each predicted exon, we have listed the genomic sequence source used for prediction. Nucleotide locations of each predicted exon are also listed. 10

15

Seq ID No: Sequence ID number
Pkey: Unique Eos probesel i
ExAccn: Exemplar Accession n Unique Eos probeset identifier number

Exemplar Accession number, Genbank accession number

UnigenelD: Unigene number
Unigene Title: Unigene gene title

20	Seq ID No:	Pkey	ExAccn	UnigeneiD	Unigene Tille
	Seq ID No: 1 & 2	410407	X66839	Hs.63287	carbonic anhydrase IX
	Seq ID No: 3 & 4	412719	AW016610	Hs.816	ESTs
25	Seq 1D No: 5 & 6	417034	NM_006183	Hs.80962	neurotensin chloride channel, calcium activated, fam
25	Seq ID No: 7 & 8	430486	BE062109	Hs.241551 Hs.38991	S100 calcium-binding protein A2
	Seq ID No: 9 & 10	407788	BE514982	Hs.38991	S100 calcium-binding protein A2
	Seq ID No: 11 & 12	407788	BE514982 BE514982	Hs.38991	S100 calcium-binding protein A2
	Seq ID No: 13 & 14	407788 407788	BE514982	Hs.38991	S100 calcium-binding protein A2
30	Seq ID No: 15 & 16 Seq ID No: 17 & 18	439285	AL133916	113.00001	hypothetical protein FLJ20093
50	Seq ID No: 19 & 20	413753	U17760	Hs.75517	taminin, beta 3 (niceln (125kD), kalinin
	Seq ID No: 21 & 22	120486	AW368377	Hs.137569	tumor protein 63 kDa with strong homolog
	Seq ID No: 23 & 24	425650		Hs.1925	desmoglein 3 (pemphigus vulgaris antigen
	Seq ID No: 25 & 26	412140	AA219691	Hs.73625	RAB6 interacting, kinesin-like (rabkines
35	Seq ID No: 27 & 28	423673	BE003054	Hs.1695	matrix metalloproteinase 12 (macrophage
	Seq ID No: 29 & 30	452838	U65011	Hs.30743	preferentially expressed antigen in mela
	Seq ID No: 31 & 32	418663	AK001100	Hs.41690	desmocollin 3
	Seq ID No: 33 & 34	418663	AK001100	Hs.41690	desmocollin 3
40	Seq ID No: 35 & 36	409632	W74001	Hs.55279	serine (or cysteine) proteinase inhibito
40	Seq ID No: 37 & 38	429610	AB024937	Hs.211092 Hs.220529	LUNX protein; PLUNC (palate lung and nas carcinoembryonic antigen-related cell ad
	Seq ID No: 39 & 40	406690 431846	M29540 BE019924	Hs.271580	uroplakin 1B
	Seq ID No: 41 & 42	418830	BE513731	Hs.88959	hypothetical protein MGC4816
	Seq ID No: 43 & 44 Seq ID No: 45 & 46	424098	AF077374	Hs.139322	small proline-rich protein 3
45	Seq ID No: 47 & 48	443648	Al085377	Hs.143610	ESTs
	Seq ID No: 49	311034	BE567130	Hs.311389	ESTs, Highly similar to NKGD_HUMAN NKG2-
	Seq ID No: 50 & 51	408522	A1541214	Hs.46320	Small proline-rich protein SPRK [human,
	Seq ID No: 52 & 53	422158	L10343	Hs.112341	protease inhibitor 3, skin-derived (SKAL
	Seq ID No: 54 & 55	435505	AF200492	Hs.211238	interleukin-1 homolog 1
50	Seq ID No: 56 & 57	417366	BE185289	Hs.1076	small proline-rich protein 1B (comifin)
	Seq ID No: 58 & 59	431958	X63629	Hs.2877	cadherin 3, type 1, P-cadherin (placenta
	Seq ID No: 60 & 61	441020	W79283	Hs.35962	ESTs
	Seq ID No: 62 & 63	423217	NM_000094	Hs.1640 Hs.11261	collagen, type VII, alpha 1 (epidermolys small proline-rich protein 2A
55	Seq ID No: 64 & 65	429538 448733	BE182592 NM_005629	Hs.187958	solute carrier family 6 (neurotransmitte
33	Seq ID No: 66 & 67 Seq ID No: 68 & 69	444371	BE540274	Hs.239	forkhead box M1
	Seq ID No: 70 & 71	444371	BE540274	Hs.239	forkhead box M1
	Seq ID No: 72 & 73	444371	BE540274	Hs.239	forkhead box M1
	Seg ID No: 74 & 75	422168	AA586894	Hs.112408	S100 calcium-binding protein A7 (psorias
60	Seq ID No: 76 & 77	422168	AA586894	Hs.112408	S100 calcium-binding protein A7 (psorias
	Seq ID No: 78 & 79	429259	AA420450	Hs.292911	Piakophilin
	Seq ID No: 80 & 81	426440	BE382756	Hs.169902	solute carrier family 2 (facilitated glu
	Seq ID No: 82 & 83	437044	AL035864	Hs.69517	differentially expressed in Fanconi's an
<i>(</i> =	Seq ID No: 84 & 85	423662	AK001035	Hs.130881	B-cell CLL/lymphoma 11A (zinc finger pro
65	Seq ID No: 86 & 87	428484	AF104032	Hs.184601 Hs.198249	solute carrier family 7 (cationic amino gap junction protein, beta 5 (connexin 3
	Seq ID No: 88 & 89	429211	AF052693 BE260964	Hs.82045	midkine (neurite growth-promoting factor
	Seq ID No: 90 & 91 Seq ID No: 92 & 93	417389 423634	AW959908	Hs.1690	heparin-binding growth factor binding pr
	Seq ID No: 94 & 95	417515	L24203	Hs.82237	ataxia-telangiectasia group D-associated
70	Seq 1D No: 96 & 97	441362	BE614410	Hs.23044	RAD51 (S. cerevisiae) homolog (E coli Re
, 0	Seg ID No: 98 & 99	425322	U63630	Hs.155637	protein kinase, DNA-activated, catalytic
	Seq ID No: 100 & 101	449003	X76342	Hs.389	alcohol dehydrogenase 7 (class IV), mu o
	Seq ID No: 102 & 103	431009	BE149762	Hs.48956	gap junction protein, beta 6 (connexin 3
	Seq ID No: 104 & 105	409103	AF251237	Hs.112208	XAGE-1 protein
75	Seq ID No: 106 & 107	417542	J04129	Hs.82269	progestagen-associated endometrial prote
	Seq ID No: 108 & 109	428471	X57348	Hs.184510	stratifin
	Seq ID No: 110 & 111	418004	U37519	Hs.87539 Hs.77256	aldehyde dehydrogenase 3 family, member enhancer of zeste (Drosophila) homolog 2
	Seq ID No: 112 & 113	414761	AU077228 X54942	Hs.83758	CDC28 protein kinase 2
80	Seq ID No: 114 & 115	418203 447343	AA256641	Hs.236894	ESTs, Highly similar to S02392 alpha-2-m
OU	Seq ID No: 116 Seq ID No: 117 & 118	437016	AU076916	Hs.5398	guanine monphosphate synthetase
	Seq ID No: 119 & 120	449230	BE613348	Hs.211579	melanoma cell adhesion molecule
	Seq ID No: 121 & 122	446989	AK001898	Hs.16740	hypothetical protein FLJ11036
	Seq ID No: 123 & 124	457819	AA057484	Hs.35406	ESTs, Highly similar to unnamed protein
85	Seq ID No: 125 & 126	424687	J05070	Hs.151738	matrix metalloproteinase 9 (gelatinase B
	*				

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	Seq ID No: 127 & 128	414430	Al346201	Hs.76118	ubiquitin carboxyl-terminal esterase L1
	Seq ID No: 129 & 130	418462	BE001596	Hs.85266	integrin, beta 4
	Seq ID No: 131 & 132	100668	L05424	Hs.169610 Hs.24763	CD44 antigen (homing function and Indian RAN binding protein 1
5	Seq ID No: 133 & 134 Seq ID No: 135 & 136	458933 418478	AI638429 U38945	Hs.1174	cyclin-dependent kinase inhibitor 2A (me
•	Seq ID No: 137 & 138	418478	U38945	Hs.1174	cyclin-dependent kinase inhibitor 2A (me
	Seq ID No: 139 & 140	418478	U38945	Hs.1174	cyclin-dependent kinase inhibitor 2A (me
	Seq ID No: 141 & 142 Seq ID No: 143 & 144	418478 446269	U38945 AW263155	Hs.1174 Hs.14559	cyclin-dependent kinase inhibitor 2A (me hypothetical protein FLJ10540
10	Seq ID No: 145 & 146	422765	AW409701	Hs.1578	baculoviral IAP repeat-containing 5 (sur
	Seq ID No: 147 & 148	436481	AA379597	Hs.5199	HSPC150 protein similar to ubiquitin-con
	Seq ID No: 149 & 150	440325 439606	NM_003812 W79123	Hs.7164 Hs.58561	a disintegrin and metalloprotelnase doma G protein-coupled receptor 87
	Seq ID No: 151 & 152 Seg ID No: 153 & 154	453884	AA355925	Hs.36232	KIAA0186 gene product
15	Seq ID No: 155 & 156	453884	AA355925	Hs.36232	KIAA0186 gene product
	Seq ID No: 157 & 158	453884	AA355925	Hs.36232	KIAA0186 gene product
	Seq ID No: 159 & 160 Seq ID No: 161 & 162	453884 404877	AA355925	Hs.36232	KIAA0186 gene product NM_005365:Homo sapiens melanoma antigen,
	Seq ID No: 163 & 164	413129	AF292100	Hs.104613	RP42 homolog
20	Seq ID No: 165 & 166	413281	AA861271	Hs.222024	transcription factor BMAL2
	Seq ID No: 167 & 168	444781	NM_014400	Hs.11950 Hs.80205	GPI-anchored metastasis-associated prote pim-2 oncogene
	Seq ID No: 169 & 170 Seq ID No: 171 & 172	416819 451320	U77735 AW118072	113.00203	dlacylgiycerol kinase, zeta (104kD)
	Seq ID No: 173 & 174	418543	NM_005329	Hs.85962	hyaluronan synthase 3
25	Seq ID No: 175 & 176	454034	NM_000691	Hs.575	aldehyde dehydrogenase 3 family, member
	Seq ID No: 177 & 178 Seq ID No: 179 & 180	425397 415817	J04088 U88967	Hs.156346 Hs.78867	topolsomerase (DNA) II alpha (170kD) protein tyrosine phosphatase, receptor-t
	Seq ID No: 181 & 182	415817	U88967	Hs.78867	protein tyrosine phosphatase, receptor-t
20	Seq ID No: 183 & 184	415817	U88967	Hs.78867	protein tyrosine phosphalase, receptor-t
30	Seq ID No: 185 & 186	415817	U88967 U88967	Hs.78867 Hs.78867	protein tyrosine phosphalase, receptor-t protein tyrosine phosphalase, receptor-t
	Seq ID No: 187 & 188 Seg ID No: 189 & 190	415817 419121	AA374372	Hs.89626	parathyrold hormone-like hormone
	Seq ID No: 191 & 192	448993	AI471630	Hs.8127	KIAA0144 gene product
25	Seq ID No: 193 & 194	421817	AF146074	Hs.108660	ATP-binding cassette, sub-family C (CFTR estrogen-responsive B box protein
35	Seq ID No: 195 & 196 Seq ID No: 197 & 198	430393 425057	BE185030 AA826434	Hs.241305 Hs.1619	achaete-scute complex (Drosophila) homol
	Seq ID No: 199 & 200	420462	AF050147	Hs.97932	chondromodulin I precursor
	Seq ID No: 201 & 202	102963	X02404	Hs.274534	calcitonin-related polypeptide, beta
40	Seq ID No: 203 & 204 Seq ID No: 205 & 206	100576 101175	X00356 U82671	Hs.37058 Hs.36980	calcitonin/calcitonin-related polypeptid melanoma antigen, family A, 2
40	Seq ID No: 207 & 208	429038	AL023513	Hs.194766	seizure related gene 6 (mouse)-like
	Seq ID No: 209 & 210	418678	NM_001327	Hs.167379	cancer/testis antigen (NY-ESO-1)
	Seq ID No: 211 & 212 Seq ID No: 213 & 214	418678 131927	NM_001327 AJ003112	Hs.167379 Hs.34780	cancer/testis antigen (NY-ESO-1) doublecortex; lissencephaly, X-linked (d
45	Seq ID No: 215 & 216	428182	BE386042	Hs.293317	ESTs, Weakly similar to GGC1_HUMAN G ANT
	Seq ID No: 217 & 218	427335	AA448542	Hs.251677	G antigen 78
	Seq ID No: 219 & 220 Seq ID No: 221 & 222	409420 114346	Z15008 AL137256	Hs.54451 Hs.130489	laminin, gamma 2 (nicein (100kD), kalini ATPase, aminophospholipid transporter-li
	Seq ID No: 223 & 224	438956	W00847	Hs.135056	Human DNA sequence from clone RP5-850E9
50	Seq ID No: 225 & 226	404440		11- 70500	NM_021048:Homo sapiens melanoma antigen,
	Seq ID No: 227 & 228 Seq ID No: 229 & 230	415669 103312	NM_005025 Y12642	Hs.78589 Hs.3185	serine (or cysteine) proteinase inhibito lysosomai
	Seq ID No: 231 & 232	320843	BE069288	Hs.34744	Homo sapiens mRNA; cDNA DKFZp547C136 (fr
E E	Seq ID No: 233	429065	A1753247	Hs.29643	Homo sapiens cDNA FLJ13103 fis, clone NT
55	Seq ID No: 234 & 235	446102 330495	AW168067 U47924	Hs.317694 Hs.71642	ESTs guanine nucleotide binding protein (G pr
	Seq ID No: 236 & 237 Seq ID No: 238	413573	AI733859	Hs.149089	ESTs
	Seq ID No: 239 & 240	428479	Y00272	Hs.334562	cell division cycle 2, G1 to S and G2 to
60	Seq ID No: 241 & 242	428479	Y00272	Hs.334562	cell division cycle 2, G1 to S and G2 to claudin 1
00	Seq ID No: 243 & 244 Seq ID No: 245	332180 437915	AF134160 Al637993	Hs.7327 Hs.202312	Homo sapiens clone N11 NTera2D1 teratoca
	Seq ID No: 246 & 247	441553	AA281219	Hs.121296	ESTs
	Seq ID No: 248 & 249	331692	A1683487	Hs.152213	wingless-type MMTV integration site fami DESC1 protein
65	Seq ID No: 250 & 251 Seq ID No: 252 & 253	429413 422283	NM_014058 AW411307	Hs.201877 Hs.114311	CDC45 (cell division cycle 45, S.cerevis
05	Seq ID No: 254 & 255	448357	N20169	Hs.108923	RAB38, member RAS oncogene family
	Seq ID No: 256 & 257	446292	AF081497	Hs.279682	Rh type C glycoprotein
	Seq ID No: 258 & 259 Seq ID No: 260 & 261	416209 453922	AA236776 AF053306	Hs.79078 Hs.36708	MAD2 (mitotic arrest deficient, yeast, h budding uninhibited by benzimidazoles 1
70	Seq ID No: 262 & 263	424046	AF027866	Hs.138202	serine (or cysteine) proteinase inhibito
	Seq ID No: 264 & 265	439223	AW238299	Hs.250618	UL16 binding protein 2
	Seq ID No: 266 & 267	429228	AI553633	Hs.326447 Hs.123114	ESTs cystatin SN
	Seq ID No: 268 & 269 Seq ID No: 270 & 271	409757 411089	NM_001898 AA456454	Hs.214291	cell division cycle 2-like 1 (PITSLRE pr
75	Seq ID No: 272 & 273	436511	AA721252	Hs.291502	ESTs
	Seq ID No: 274 & 275	428969	AF120274	Hs.194689	artemin
	Seq ID No: 276 & 277 Seq ID No: 278 & 279	428969 428969	AF120274 AF120274	Hs.194689 Hs.194689	artemin artemin
	Seq ID No: 280 & 281	428969	AF120274	Hs.194689	artemin
80	Seq ID No: 282	407137	T97307	Un 225054	gb:ye53h05.s1 Soares fetal liver spieen
	Seq ID No: 283 & 284 Seq ID No: 285 & 286	412723 450701	AA648459 H39960	Hs.335951 Hs.288467	hypothetical protein AF301222 hypothetical protein XP_098151 (leucine-
	Seq ID No: 287 & 288	405770	,		NM_002362:Homo sapiens melanoma antigen,
0.5	Seq ID No: 289 & 290	439453	BE264974	Hs.6566	thyroid hormone receptor interactor 13
85	Seq ID No: 291 & 292	414774	X02419	Hs.77274	plasminogen activator, urokinase

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	Seq ID No: 293 & 294	424629	M90656	Hs.151393	glutamate-cysteine ligase, catalytic sub
	Seq ID No: 295 & 296	437789	Al581344	Hs.127812	ESTs, Weakly similar to T17330 hypotheti
	Seq ID No: 297 & 298	437789	Al581344	Hs.127812	ESTs, Weakly similar to T17330 hypotheti
5	Seq ID No: 299 & 300	437789	Al581344	Hs.127812 Hs.127812	ESTs, Weakly similar to T17330 hypotheti ESTs, Weakly similar to T17330 hypotheti
,	Seq ID No: 301 & 302 Seq ID No: 303 & 304	437789 437789	AI581344 AI581344	Hs.127812	ESTs, Weakly similar to T17330 hypotheti
	Seq ID No: 305 & 306	453968	AA847843	Hs.62711	High mobility group (nonhistone chromoso
	Seq ID No: 307 & 308	403478			NM_022342:Homo sapiens kinesin protein 9
10	Seq ID No: 309	441525	AW241867	Hs.127728	ESTs
10	Seq ID No: 310 & 311 Seq ID No: 312 & 313	434105 428810	AW952124 AF068236	Hs.13094 Hs.193788	presenilins associated rhombold-like pro nitric oxide synthase 2A (inducible, hep
	Seq ID No: 314 & 315	413691	AB023173	Hs.75478	ATPase, Class VI, type 11B
	Seq ID No: 316 & 317	423934	U89995	Hs.159234	forkhead box E1 (thyroid transcription f
15	Seq ID No: 318 & 319	409228	R16811	Hs.22010	ESTs, Weakly similar to 2109260A B cell
13	Seq ID No: 320 & 321 Seq ID No: 322 & 323	425734 413582	AF056209 AW295647	Hs.159396 Hs.71331	peptidylglycine alpha-amidating monooxyg hypothetical protein MGC5350
	Seq ID No: 324 & 325	438403	AA806607	Hs.292206	ESTs
	Seq ID No: 326 & 327	403329			unnamed protein product [Homo sapiens]
20	Seq ID No: 328 & 329	409893	AW247090	Hs.57101	minichromosome maintenance deficient (S.
20	Seq ID No: 330 & 331 Seq ID No: 332 & 333	119073 113195	BE245360 H83265	Hs.279477 Hs.8881	v-ets erythroblastosis virus E26 oncogen ESTs, Weakly similar to S41044 chromosom
	Seq ID No: 334 & 335	102283	AW161552	Hs.83381	guanine nucleotide binding protein 11
	Seq ID No: 336 & 337	101345	NM_005795	Hs.152175	calcitonin receptor-like
25	Seq ID No: 338 & 339	103280	U84722	Hs.76206	cadherin 5, type 2, VE-cadherin (vascula
25	Seq ID No: 340 & 341 Seq ID No: 342 & 343	102012 105729	BE259035 H46612	Hs.118400 Hs.293815	singed (Drosophila)-like (sea urchin fas Homo sapiens HSPC285 mRNA, partial cds
	Seq ID No: 344 & 345	134299	AW580939	Hs.97199	complement component C1q receptor
	Seq ID No: 346 & 347	412719	AW016610	Hs.816	ESTs
20	Seq ID No: 348 & 349	422158	L10343	Hs.112341	protease Inhibitor 3, skin-derived (SKAL
30	Seq ID No: 350 & 351 Seq ID No: 352 & 353	128924 100486	BE279383 T19006	Hs.26557 Hs.10842	plakophilin 3 RAN, member RAS oncogene family
	Seq ID No: 354 & 355	419121	AA374372	Hs.89626	parathyroid hormone-like hormone
	Seq ID No: 356 & 357	409459	D86407	Hs.54481	low density lipoprotein receptor-related
35	Seq ID No: 358 & 359	330493	M27826	Us 92772	endogenous retroviral protease
33	Seq ID No: 360 & 361 Seq ID No: 362 & 363	417866 418113	AW067903 Al272141	Hs.82772 Hs.83484	collagen, type XI, alpha 1 SRY (sex determining region Y)-box 4
	Seq ID No: 364 & 365	437016	AU076916	Hs.5398	guanine monphosphate synthetase
	Seq ID No: 366 & 367	429612	AF062649	Hs.252587	pituitary tumor-transforming 1
40	Seq ID No: 368 & 369	440704	M69241	Hs.162 Hs.286145	insulin-like growth factor binding prote SRB7 (suppressor of RNA polymerase B, ye
40	Seq ID No: 370 & 371 Seq ID No: 372 & 373	431221 431565	AA449015 AF161470	Hs.260622	butyrate-Induced transcript 1
	Seq ID No: 374 & 375	431565	AF161470	Hs.260622	butyrate-induced transcript 1
	Seq ID No: 376 & 377	132354	BE185289	Hs.1076	small proline-rich protein 18 (comifin)
45	Seq ID No: 378 & 379	424441 103768	X14850 AF086009	Hs.147097 Hs.296398	H2A histone family, member X gb:Homo sapiens full length insert cDNA
73	Seq ID No: 380 & 381 Seq ID No: 382 & 383	417512	X76534	Hs.82226	glycoprotein (transmembrane) nmb
	Seq ID No: 384 & 385	425266	J00077	Hs.155421	alpha-fetoprotein
	Seq ID No: 386 & 387	424503	NM_002205	Hs.149609	integrin, alpha 5 (fibronectin receptor, matrix metalloproteinase 10 (stromelysin
50	Seq ID No: 388 & 389 Seq ID No: 390 & 391	400289 418007	X07820 M13509	Hs.2258 Hs.83169	matrix metalloproteinase 1 (interstitial
50	Seq ID No: 392 & 393	418007	M13509	Hs.83169	matrix metalloproteinase 1 (interstitial
	Seq ID No: 394 & 395	418738	AW388633	Hs.6682	solute carrier family 7, (cationic amino
	Seq ID No: 396 & 397	415138	C18356 AA084248	Hs.295944 Hs.85339	tlssue factor pathway inhibitor 2 G protein-coupled receptor 39
55	Seq ID No: 398 & 399 Seq ID No: 400 & 401	418506 423961	D13666	Hs.136348	periostin (OSF-20s)
	Seq ID No: 402 & 403	414812	X72755	Hs.77367	monokine induced by gamma interferon
	Seq ID No: 404 & 405	417433	BE270266	Hs.82128	5T4 oncofetal trophoblast glycoprotein -
	Seq ID No: 406 & 407 Seq ID No: 408 & 409	417433 422867	BE270266 L32137	Hs.82128 Hs.1584	5T4 oncofetal trophoblast glycoprotein cartilage oligomeric matrix protein (pse
60	Seq ID No: 410 & 411	428227	AA321649	Hs.2248	small inducible cytokine subfamily 8 (Cy
	Seq ID No: 412 & 413	444381	BE387335	Hs.283713	ESTs, Weakly similar to S64054 hypotheti
	Seq ID No: 414 & 415	400303 411789	AA242758 AF245505	Hs.79136 Hs.72167	LIV-1 protein, estrogen regulated Adlican
	Seq ID No: 416 & 417 Seq ID No: 418 & 419	428698	AA852773	Hs.334838	KIAA1866 protein
65	Seq ID No: 420 & 421	450098	W27249	Hs.8109	hypothetical protein FLJ21080
	Seq ID No: 422 & 423	421552	AF026692	Hs.105700	secreted frizzled-related protein 4
	Seq ID No: 424 & 425 Seq ID No: 426 & 427	452747 450375	BE153855 AA009647	Hs.61460	lg superfamily receptor LNIR a disintegrin and metalloproteinase doma
	Seq ID No: 428 & 429	426215	AW963419	Hs.155223	stanniocalcin 2
70	Seq ID No: 430 & 431	425247	NM_005940	Hs.155324	matrix metalloproteinase 11 (stromelysin
	Seq ID No: 432 & 433	432201	AI538613	Hs.298241	Transmembrane protease, serine 3 collagen, type X, alpha 1 (Schmid metaph
	Seq ID No: 434 & 435 Seq ID No: 436 & 437	427585 442117	D31152 AW664964	Hs.179729 Hs.128899	ESTs; hypothetical protein for IMAGE:447
	Seq ID No: 438 & 439	431211	M86849	Hs.323733	gap junction protein, beta 2, 26kD (conn
75	Seq ID No: 440 & 441	447033	A1357412	Hs.157601	ESTs
	Seq ID No: 442 & 443	447033	AI357412	Hs.157601	ESTs ESTs
	Seq ID No: 444 & 445 Seq ID No: 446 & 447	447033 115522	AI357412 BE614387	Hs.157601 Hs.333893	c-Myc target JPO1
	Seq ID No: 448 & 449	410418	D31382	Hs.63325	transmembrane protease, serine 4
80	Seq ID No: 450 & 451	409041	AB033025	Hs.50081	Hypothetical protein, XP_051860 (KIAA119
	Seq ID No: 452 & 453	409041	AB033025	Hs.50081 Hs.108106	Hypothetical protein, XP_051860 (KIAA119 transcription factor
	Seq ID No: 454 & 455 Seq ID No: 456 & 457	452461 412420	N78223 AL035668	Hs.73853	bone morphogenetic protein 2
0.7	Seq ID No: 458 & 459	416658	U03272	Hs.79432	fibrillin 2 (congenital contractural ara
85	Seq ID No: 460 & 461	407811	AW190902	Hs.40098	cysteine knot superfamily 1, BMP antagon

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	Seq ID No: 462 & 463	437852	BE001836	Hs.256897	ESTs, Weakly similar to dJ365O12.1 [H.sa
	Seq ID No: 464 & 465	402075			ENSP00000251056*:Plasma membrane calcium
	Seq ID No: 466 & 467	421110	AJ250717	Hs.1355	cathepsin E
	Seq ID No: 468 & 469	451668	Z43948	Hs.326444	cartilage acidic protein 1
5	Seq ID No: 470 & 471	451668	Z43948	Hs.326444	cartilage acidic protein 1
•	Seq ID No: 472 & 473	451668	Z43948	Hs.326444	cartilage acidic protein 1
	Seq ID No: 474 & 475	422282	AF019225	Hs.114309	apolipoprotein L
	Seq ID No: 476 & 477	425852	AK001504	Hs.159651	death receptor 6, TNF superfamily member
	Seq ID No: 478 & 479	439738	BE246502	Hs.9598	sema domain, immunoglobulin domain (ig),
10	Seq ID No: 480 & 481	427747	AW411425	Hs.180655	serine/threonine kinase 12
10	Seq ID No: 482 & 483	420281	Al623693	Hs.323494	Predicted cation efflux pump
		405932	71020000	110.020101	C15000305:gi 3806122 gb AAC69198.1 (AF0
	Seq ID No: 484 & 485	405932			C15000305:glj3806122[gb]AAC69198.1] (AF0
	Seq ID No: 486 & 487	444342	NM_014398	Hs.10887	similar to lysosome-associated membrane
15	Seq ID No: 488 & 489	421379	Y15221	Hs.103982	small inducible cytokine subfamily B (Cy
13	Seq JD No: 490 & 491	421379	U65590	Hs.81134	Interleukin 1 receptor antagonist
	Seq ID No: 492 & 493	430890	X54232	Hs.2699	glypican 1
	Seq ID No: 494 & 495			Hs.288650	aquaporin 4
	Seq ID No: 496 & 497	419721	NM_001650 AB020684	Hs.11217	KIAA0877 protein
20	Seq ID No: 498 & 499	444471		Hs.75184	chitinase 3-like 1 (cartilage glycoprote
20	Seq ID No: 500 & 501	413063	AL035737 Al034361	Hs.135150	lung type-i cell membrane-associated gly
	Seq ID No: 502 & 503	433800		Hs.29352	tumor necrosis factor, alpha-induced pro
	Seq ID No: 504 & 505	452401	NM_007115	Hs.29352	tumor necrosis factor, alpha-induced pro
	Seq ID No: 506 & 507	452401	NM_007115		solute carrier family 6 (neurotransmitte
25	Seq ID No: 508 & 509	450001	NM_001044	Hs.406 Hs.63287	carbonic anhydrase IX
25	Seq ID No: 510 & 511	410407	X66839	H8.03207	gb:hd13d01.x1 Soares_NFL_T_GBC_S1 Homo s
	Seq ID No: 512 & 513	309931	AW341683	Un 046	ESTs
	Seq ID No: 514 & 515	412719	AW016610	Hs.816	
	Seq ID No: 516 & 517	417034	NM_006183	Hs.80962	neurotensin chloride channel, calcium activated, fam
20	Seq ID No: 518 & 519	430486	BE062109	Hs.241551	laminin, beta 3 (nicein (125kD), kalinin
30	Seq ID No: 520 & 521	413753	U17760	Hs.75517	desmoglein 3 (pemphigus vulgaris antigen
	Seq ID No: 522 & 523	425650	NM_001944	Hs.1925	matrix metalloproteinase 12 (macrophage
	Seq ID No: 524 & 525	423673	BE003054	Hs.1695	desmocallin 3
	Seq ID No: 526 & 527	418663	AK001100	Hs.41690	
25	Seq ID No: 528 & 529	418663	AK001100	Hs.41690	desmocollin 3 LUNX protein; PLUNC (palate lung and nas
35	Seq ID No: 530 & 531	429610	AB024937	Hs.211092	carcinoembryonic antigen-related cell ad
	Seq ID No: 532 & 533	406690	M29540	Hs.220529	
	Seq ID No: 534 & 535	431846	BE019924	Hs.271580	uroplakin 1B protease inhibitor 3, skin-derived (SKAL
	Seq ID No: 536 & 537	422158	L10343	Hs.112341	cadherin 3, type 1, P-cadherin (placenta
40	Seq ID No: 538 & 539	431958	X63629	Hs.2877	differentially expressed in Fanconi's an
40	Seq ID No: 540 & 541	437044	AL035864	Hs.69517	solute carrier family 7 (cationic amino
	Seq ID No: 542 & 543	428484	AF104032	Hs.184601	gap junction protein, bela 5 (connexin 3
	Seq ID No: 544 & 545	429211	AF052693	Hs.198249	midkine (neurite growth-promoting factor
	Seq ID No: 546 & 547	417389	BE260964	Hs.82045	gap junction protein, beta 6 (connexin 3
45	Seq ID No: 548 & 549	431009	BE149762	Hs.48956	progestagen-associated endometrial prote
43	Seq ID No: 550 & 551	417542	J04129	Hs.82269	melanoma cell adhesion molecule
	Seq ID No: 552 & 553	449230	BE613348	Hs.211579	a disintegrin and metalloproteinase doma
	Seq ID No: 554 & 555	410555	U92649	Hs.64311	a disintegrin and metalloproteinase doma
	Seq ID No: 556 & 557	410555	U92649	Hs.64311	matrix metalloproteinase 9 (gelatinase B
50	Seq ID No: 558 & 559	424687	J05070	Hs.151738 Hs.85266	Integrin, beta 4
50	Seq ID No: 560 & 561	418462	BE001596 AA381807	Hs.61762	hypoxia-inducible protein 2
	Seq ID No: 562 & 563	410274 439606	W79123	Hs.58561	G protein-coupled receptor 87
	Seq ID No: 564 & 565	404877	VV/5123	113.50501	NM 005365:Homo sapiens melanoma antigen,
	Seq ID No: 566 & 567 Seq ID No: 568 & 569	444781	NM_014400	Hs.11950	GPI-enchored metastasis-associated prote
55		418543	NM_005329	Hs.85962	hyaturonan synthase 3
55	Seq ID No: 570 & 571 Seq ID No: 572 & 573	415817	U88967	Hs.78867	protein tyrosine phosphatase, receptor-t
	Seq ID No: 574 & 575	415817	U88967	Hs.78867	protein tyrosine phosphatase, receptor-t
	Seq ID No: 576 & 577	415817	U88967	Hs.78867	protein tyrosine phosphatase, receptor-t
	Seq ID No: 578 & 579	415817	U88967	Hs.78867	protein tyrosine phosphatase, receptor-t
60	Seq ID No: 580 & 581	415817	U88967	Hs.78867	protein tyrosine phosphalase, receptor-t
-	Seq ID No: 582 & 583	415817	U88967	Hs.78867	protein tyrosine phosphatase, receptor-t
•	Seq ID No: 584 & 585	421817	AF146074	Hs.108660	ATP-binding cassette, sub-family C (CFTR
	Seg ID No: 586 & 587	418678	NM_001327	Hs.167379	cancer/testis antigen (NY-ESO-1)
	Seq ID No: 588 & 589	418678	NM_001327	Hs.167379	cancer/testis antigen (NY-ESO-1)
65	Seq ID No: 590 & 591	409420	Z15008	Hs.54451	laminin, gamma 2 (nicein (100kD), kalini
00	Seq ID No: 592 & 593	332180	AF134160	Hs.7327	claudin 1
	Seq ID No: 594 & 595	408790	AW580227	Hs.47860	neurotrophic tyrosine kinase, receptor,
	Seq ID No: 596 & 597	408790	AW580227	Hs.47860	neurotrophic tyrosine kinase, receptor,
	Seg ID No: 598 & 599	439223	AW238299	Hs.250618	UL16 binding protein 2
70	Seq ID No: 600 & 601	409757	NM_001898	Hs.123114	cystatin SN
	Seq ID No: 602 & 603	428969	AF120274	Hs.194689	artemin
	Seq ID No: 604 & 605	428969	AF120274	Hs.194689	artemin
	Seq ID No: 606 & 607	428969	AF120274	Hs.194689	artemin
	Seq ID No: 608 & 609	428969	AF120274	Hs.194689	artemin
75	Seq ID No: 610 & 611	450701	H39960	Hs.288467	hypothetical protein XP_098151 (leucine-
	Seg ID No: 612 & 613	450701	H39960	Hs.288467	hypothetical protein XP_098151 (leucine-
	Seq ID No: 614 & 615	414774	X02419	Hs.77274	plasminogen activator, urokinase
	Seq ID No: 616 & 617	407944	R34008	Hs.239727	desmocolin 2
	Seq ID No: 618 & 619	407944	R34008	Hs.239727	desmocollin 2
80	Seq ID No: 620 & 621	457489	Al693815	Hs.127179	cryptic gene
	Seq ID No: 622 & 623	429547	AW009166	Hs.99376	ESTs
	Seq ID No: 624 & 625	407242	M18728	•	gb:Human nonspecific crossreacting antig
	Seq ID No: 626 & 627	407242	M18728		gb:Human nonspecific crossreacting antig
~-	Seq ID No: 628 & 629	407242	M18728		gb:Human nonspecific crossreacting antig
85	Seq ID No: 630 & 631	444006	BE395085	Hs.10086	type I transmembrane protein Fn14

PCT/US02/12476 WO 02/086443 Seq ID No: 632 & 633 NM_003816 Hs.2442 a disintegrin and metalloproteinase doma Hs.1473 Hs.288433 Seq ID No: 634 & 635 422109 S73265 gastrin-releasing peptide Seq ID No: 636 & 637 Seq ID No: 638 & 639 AW470411 419235 neurotrimin Hs.22920 similar to S68401 (cattle) glucose induc 449048 Z45051 5 Seq ID No: 640 & 641 419216 AU076718 Hs.164021 small inducible cytokine subfamily B (Cy granin-like neuroendocrine peptide precu Seq ID No: 642 & 643 431462 AW583672 Hs.256311 Hs.52620 Seq ID No: 644 & 645 448243 AW369771 integrin, heta 8 Seq ID No: 646 & 647 Seq ID No: 648 & 649 Hs.169840 TTK protein kinase 426427 M86699 AJ245671 Hs.12844 EGF-like-domain, multiple 6 445537 10 Seq ID No: 650 & 651 422278 AF072873 Hs.114218 frizzled (Drosophila) homolog 6 Seq ID No: 652 & 653 Seq ID No: 654 & 655 Seq ID No: 656 & 657 Seq ID No: 658 & 659 KIAA0175 gene product secreted phosphoprotein 1 (osteopontin, NM_014791 AU076643 Hs.184339 428450 446619 Hs.313 Hs.32964 SRY (sex determining region Y)-box 11 453392 U23752 426514 BE616633 Hs.170195 bone morphogenetic protein 7 (osteogenic parathyroid hormone receptor 2 15 Seq ID No: 660 & 661 425776 U25128 Hs.159499 parathyroid hormone receptor 2 endothelial differentiation, lysophospha Seq ID No: 662 & 663 Seq ID No: 664 & 665 Seq ID No: 666 & 667 Seq ID No: 668 & 667 Hs. 159499 425776 431515 1125128 NM_012152 Hs.258583 PTK7 protein tyrosine kinase 7 ESTs, Weakly similar to JC7328 amino aci ESTs, Weakly similar to JC7328 amino aci Hs.90572 419452 U33635 432653 N62096 Hs.293185 20 Seq ID No: 670 & 671 Seq ID No: 672 & 673 Seq ID No: 674 & 675 Hs.293185 N62096 432653 N62096 Hs.293185 ESTs, Weakly similar to JC7328 amino aci 432653 432653 N62096 Hs.293185 ESTs, Wealdy similar to JC7328 amino aci Seq ID No: 676 & 677 410001 AB041036 AW043782 Hs.57771 Hs.293616 kallikrein 11 Seq ID No: 678 & 679 Seq ID No: 680 & 681 Seq ID No: 682 & 683 **ESTs** 426501 25 408369 R38438 Hs.182575 solute carrier family 15 (H??? transport CGI-147 protein prostate differentiation factor 445413 AA151342 Hs.12677 Hs.296638 Seq ID No: 684 & 685 422424 Al186431 matrix metalloproteinase 7 (matrilysin, Seq ID No: 686 & 687 Seq ID No: 688 & 689 428330 1 22524 Hs 2256 distal-less homeo box 5 420610 AI683183 Hs.99348 30 TABLE 15B Unique Eos probeset identifier number Pkey: Unique Eos probesel CAT number: Gene d'uster number 35 Genbank accession numbers CAT Number Accession 309931 AW341683 M27826 R78416 AA307645 AW957879 AW957800 AA633529 H03662
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AW118072 Al631982 T15734 AA224195 Al701458 W20198 F26326 AA890570 N90552 AW071907 Al671352 Al375892 T03517 R88265
Al124088 AA224388 Al084316 Al354686 T33652 Al140719 Al720211 T03490 Al372637 T15415 AW205836 AA630384 T03515 T33230 330493 33264_5 40 439285 47065 1 450375 83327_1 86576_1 451320 45 AA017131 AA443303 T33623 AI222556 T33511 T33785 AI419606 D55612 TABLE 15C 50 Unique number corresponding to an Eos probeset Sequence source. The 7 digit numbers in this column are Genbank Identifier (GI) numbers. "Dunham I. et al." refers to the publication entitled "The DNA Ref: sequence of human chromosome 22." Dunham I. et al., Nature (1999) 402:489-495. Indicates DNA strand from which exons were predicted. 55 Nt_position: Indicates nucleotide positions of predicted exons. Nt_position 121907-122035,122804-122921,124019-124161,124455-124610,125672-126076 Pkey 8117407 402075 Plus 96450-96598 8516120 403329 Plus 60 116458-116564 403478 9958258 Plus 404440 7528051 80430-81581 1095-2107 404877 1519284 Plus

61057-62075

123525-123713

2735037

7767812

Phis

Minus

405770

405932

65

WO 02/086443

Table 16

5

Seq ID NO: 1 DNA sequence Nucleic Acid Accession #: NM_001216 Coding sequence: 43..1422

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PCT/US02/12476

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Seq ID NO: 10 Protein sequence:

Protein Accession #: NP_005969.1

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PCT/US02/12476

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8 <i>5</i>		35 DNA sequ					
		ld Accession		quence			
	coging sequ	lence: 146-	1273				

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21
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Seq ID NO: 38 Protein sequence: Protein Accession #: NP_057667

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51

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Nucleic Acid Accession #: Eos sequence

PCT/US02/12476

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	Nucleic Actions sequently	Id Accession uence: 99-89 II GEGEGTEGGE TCCTAGCTGA CGCCGGGATC CACCGCCTTTC CAGCAATTTC AGCCAGTGCA	#: NM_0000 21 TCGGACCTGC CGGCTTTTAC CTGGCAGAGG TACGCCGCTG TACGCAGGTCC CAGGGTGTGC	31 CAAGGCCACC TGCCTAGGAT CGCCCCGAGT ACATTGTGT GCAGCTTTCT GCTTTGCCAC	GCAGGGGGA GACGCTGCGG GCGAGCCCAG CTTACTGGAT CGAAGGGCTG AGTGCAGTAC	GCAAGGGACA CTTCTGGTGG CACAGGGAGA GGCTCCTCAT GTGCTGCCTT AGCGATGACC	120 180 240 300 360
50	Nucleic According sequences of the control of the c	Id Accession uence: 99-89 11 GGCGCTGGGC TCCTAGCTGA CGCCGGGATC CACGCGCCTT CAGCAATTTC AGCCAGTGCA GTTCGGCCTG	#: NM_0000 21 TCGGACCTGC CGGCTTTTAC CTGCAGAGG TACGCCGCTG CGCGAGGTCC CAGGGTTGC CAGGGTTGC CAGGGTTGC CAGGGTTGC	31 CAAGGCCACC TGCCTAGGAT CGCCCGAGT ACATTGTGTT GCAGCTTTCCT GCTTTGCCAG GCTCTGGGGG	GCAGGGGGA GACGCTGCGG GCGAGCCCAG CTTACTGGAT CGAAGGGCTG AGTGCAGTAC TGATGTGATC	GCAAGGGACA CTTCTGGTGG CACAGGGAGA GGCTCCTCAT GTGCTGCCTT AGCGATGACC CGCGCCATCC	120 180 240 300 360 420
50	Nucleic Actions sequently	Id Accession uence: 99-8: II GGCGCTGGGC TCCTAGCTGA CGCCGGGATT CAGCAGTTTC AGCAATTTC AGCCAGTGCA GTTCGGCCTG CTACAAGGGG	#: NM_0000 21 TGGGACCTGC CGGCTTTTAC CTGGCAGAGG TACGCCGCTG CGCAGGTCC CAGGGTGTGC CAGGGTGTGC GGCAACACTC	31 CAAGGCCACC TGCCTAGGAT CGCCCCGAGT ACATTGTGTT GCAGCTTTCT GCTTTGGCAC GCTCTGGGGG GCACAGGGGC	GCAGGGGGA GACGCTGCGG GCGAGCCCAG CTTACTGGAT CGAAGGGCTG AGTGCAGTAC TGATGTGATC TGCAATTCTC	GCAAGGGACA CTTCTGGTGG CACAGGGAGA GGCTCCTCAT GTGCTGCCTT AGCGATGACC CGCGCCATCC CATGTGGCTG	120 180 240 300 360 420 480
50	Nucleic According sequences of the control of the c	Id Accession nence: 99-8: II GGCGCTGGGC TCCTAGCTGA CGCCGGATTC CACGGCCTT CAGCAATTTC AGCCAGTGCA GTTCGGCCTG CTACAAGGGG CCTGCCCCAG	#: NM_0000 21 TCGGACCTGC CGGCTTTTAC CTGGCAGAGG TACGCCGCTG CGCAGGTGCC CAGGGTGTGC CAGGGTGTGC CAGGGTGTGC CAGGGTGTGC CAGGGTGTGC CAGGGTGTGC CTGGCCCGAC	31 CAAGGCCACC TGCCTAGGAT CGCCCCGAGT ACATTGTTT GCTTTGCCAC GCTCTGGGGG GCTCTGGGGG CTGGTGTCCC	GCAGGGGGA GACGCTGCGG GCGAGCCCAG CTTACTGGAT CGAAGGGCTG AGTGCAGTAC TGATGTGATC TGCAATTCTC CAAGGTCTGC	GCAAGGGACA CTTCTGGTGG CACAGGGAGA GGCTCCTCAT GTGCTGCCTT AGCGATGACC CGCGCCATCC CATGTGGCTG ATCCTGATCA	120 180 240 300 360 420 480 540
50 55	Nucleic According sequence of the control of the co	Id Accession uence: 99-89 11 GGCGCTGGGC TCCTAGCTGA CGCCGGGATC CACGCGCTTT CAGCAATTTC AGCCAGTGCA GTTCGGCCTG CTACAAGGGG CTCCAGGAGC GTCCCAGGAC	#: NM_0000 21 TCGGACCTGC CGGCTTTTAC CTGGCAGAGG TACGCCGCTG CAGGGTGTGC CAGGGTGTGC CAGGTGTGC CTGGCCCGAC CTGGTGGACAC	31 CAAGGCCACC TGCCTAGGAT CGCCCCGAGT ACATTGTGT GCTTTGCCAC GCTCTGGGGG GCACAGGGGC CTGGTGTCCC CAGCTGCCCA	CAGGGGGGA GACGCTGCGG GCGAGCCCAG CTTACTGGAT CGAAGGGCTG AGTGCAGTAC TGATGTGATC TGCAATTCTC CAAGGTCTGC AAGGCTGAAG	GCAAGGGACA CTTCTGGTGG CACAGGGAGA GGCTCCTCAT GTGCTGCCTT AGCGATGACC CGCGCCATCC CATGTGGCTG ATCCTGATCA GGGCAGGGGG	120 180 240 300 360 420 480
50	Nucleic According sequence of the control of the co	Id Accession Lence: 99-8: II GGCGCTGGGC TCCTAGCTGA CGCGGGATC CACGGCCTT CAGCAATTTC AGCAGTGCA GTTCGGCTG CTACAAGGG CCTGCCCAG GTCCCAGGAC TGCTGTGGGG TCCCAGGAC TGCTGTGGGG	#: NM_0000 21 TCGGACCTGC CGGCTTTTAC CTGGCAGAGG TACGCCGCTG CGCAGGTGCC CAGGGTGTGC CAGGGTGTGC CAGGGTGTGC CTGGCCGAC CTGGCCCGAC CTGGCCGAC ATCAAGAATG	31 CAAGGCCACC TGCCTAGGAT CGCCCCGAGT ACATTGTTT GCTTTGCCAC GCTCTGGGGG GCACAGGGGC CTGGTGTCCC CAGCTGCCCA CTGACCTTGA	CAGGGGGGA GAGGCCAG GCGAGCCAG GCGAGCCAG CTTACTGGAT CGAAGGGCTG AGTGCAGTAC TGCAATTCTC CAAGGTCTGAAG GAGCTGAAG GAGCTGAAG	GCAAGGGACA CTTCTGGTGG CACAGGGAGA GGCTCCTCAT GTGCTGCCTT AGCGATGACC CAGGGCATCC CATGTGGCTG ATCCTGATCA GGCAGGGGG GGAGTTGCCT AGGACACTAC	120 180 240 300 360 420 480 540 600
50 55	Nucleic According sequence of the control of the co	Id Accession Lence: 99-8: II GGCGCTGGGC TCCTAGCTGA CGCGGGATC CACGGCCTT CAGCAATTTC AGCAGTGCA GTTCGGCTG CTACAAGGG CCTGCCCAG GTCCCAGGAC TGCTGTGGGG TCCCAGGAC TGCTGTGGGG	#: NM_0000 21 TCGGACCTGC CGGCTTTTAC CTGGCAGAGG TACGCCGCTG CGCAGGTGCC CAGGGTGTGC CAGGGTGTGC CAGGGTGTGC CTGGCCGAC CTGGCCCGAC CTGGCCGAC ATCAAGAATG	31 CAAGGCCACC TGCCTAGGAT CGCCCCGAGT ACATTGTTT GCTTTGCCAC GCTCTGGGGG GCACAGGGGC CTGGTGTCCC CAGCTGCCCA CTGACCTTGA	CAGGGGGGA GAGGCCAG GCGAGCCAG GCGAGCCAG CTTACTGGAT CGAAGGGCTG AGTGCAGTAC TGCAATTCTC CAAGGTCTGAAG GAGCTGAAG GAGCTGAAG	GCAAGGGACA CTTCTGGTGG CACAGGGAGA GGCTCCTCAT GTGCTGCCTT AGCGATGACC CAGGGCATCC CATGTGGCTG ATCCTGATCA GGCAGGGGG GGAGTTGCCT AGGACACTAC	120 180 240 300 360 420 480 540 600 660
50 55	Nucleic Act Coding sequence	Id Accession uence: 99-8: 11 GGGGCTGGGC TCCTAGCTGA CGCCGGGATT CAGCAATTTC AGCCAGTGCA AGTTCGCCTG CTACAAGGG CCTGCCCAG GTCCCAGGAC TCCTGCGACT TCCGACTTC TTCCCGAGAG ACCTCTGCT	1 #: NM_0000 333 21 TGGGACCTGC CGGCTTTTAC CTGGCAGAGG TACGCCGCTG CGCAGGTCTC CAGGGTGTGC CAGGGTGTGC CTGGCCCGAC CTGGTGGACA ATCAAGAATG TTCTTCTTCG GTGTGCACGA CCACGAGACCC CTGGTGCACAA	31 CAAGGCCACC TGCCTAGGAT CGCCCCGAGT ACATTGTGT GCAGCTTTCT GCTCTGGGGG GCACAGGGGC CTGGTGTCCC CAGCTGCCCA CTGACCCTGA TCAATGACTT CTGCTGGTG TGTGTCT CTGCTGGTGT	GCAGGGGGA GCAGGGGGGA GCAAGCCCAG CTTACTGGAT CGAAGGGCTA TGATGTGAT TGATGTGAT CCAAGTCTGC AAGGCTGAAG GGAGCTGAAG CAGCATCTTG TGAGCATCTTG TGAGCCAAGC	GCAAGGGACA CTTCTGGTGG CACAGGGAGA GGCTCCTCAT GTGCTGCCTT AGCGATGACC CACGGCCATCC CATGTGGCTG ATCCTGATCA GGGCAGGGGG CGAGTTGCCT AGGACACTAC ACCCGACCTC AGCCAATCCT	120 180 240 300 360 420 480 540 600 660 720
50 55 60	Nucleic According sequences of the control of the c	Id Accession uence: 99-8: II GGCGCTGGGC TCCTAGCTGA CGCGGGATTC CACGGGCTT AGCAATTTC AGCAGTGCA GTTCGGCTG GTCCCAGGAC TCCCAGGAC TCCCAGGAC TCCCAGGAC TCCCGGACTTC TTCCCGGAGA GACCTCTGCT	#: NM_0000 21 TGGGACCTGC CGGCTTTTAC CTGGCAGAGG TACGCCGCTG CGGAGGTCC CAGGGTGTGC GATGCACTTC CTGGCCGAC CTGGTGGACA ATCAAGAATG TTCTTCTCG GTGTGACAC GTGTGACAC GTGTGACAC CTGGTGACAC CTGGTGACAC CTGGTGACAC CTGGTGACAC CTGGTGACAC CTGGTGACAC CTGGTGACAC CTGGTGACAC CTGGTGCACGAC CGCACGAGACC CCACGAGACC CCCAGTGGCC	31 CAAGGCCACC TGCCTAGGAT CGCCCGAGT ACATTGTTT GCTTTGCCAC GCTCTGGGGG CCAGGGGGC CTGGTGTCCC AGCTGCCCA CTGACCCTGA TCAATGACTT CTGCTGGTGT CTGTGGTGT CTGTGGTGT CTGTGGTGT	CAGAGGGGGA GAGAGCCAG GCGAGCCAG GCGAGCCAG GCGAGCCAG AGTGCAGTAC TGATGTGATC CAGAGTCTGC AAGGCTGAAG GGAGCTGAAG CAGCATCTTG CAGACTCTTG CAGACTCTGC TGAAGGCTAAGC CTACAAGGTC	GCAAGGGACA CTTCTGGTGG CACAGGGAGA GGCTCCTCAT GGCGCATCC CAGGCCATCC CATGTGGCTG ATCCTGATCA GGCAGGGGG CGAGTTGCCT AGGACACTAC ACCGACCTC CAGCAATCCT CAGTACACTC	120 180 240 300 360 420 480 540 600 720 780 840 900
50 55	Nucleic According sequences of the control of the c	Id Accession lence: 99-8: II GGCGCTGGGC TCCTAGCTGA CGCGGGATC CACGGCCTT AGCAATTTC AGCAGTGCA GTTCGGCTGA GTCCCAGG CTGCCCAG GTCCCAGGAC TCCTGTGGGG CTCCGACTTC TTCCCGAGAA GACCTCTGCT GTGGACAGC GTGGACAGC GTGGACAGC GTGGACAGC	#: NM_0000 21 TCGGACCTGC CGGCTTTTAC CTGGCAGAGG TACGCCGCTG GAGGGTCC CAGGGTGTGC GATGCACATC CTGGCCGAC CTGGTGGACA ATCAAGAATG TTCTTCG GTGTGCACGA CCACGAGACC CGCAGTGCC CCCACTGCCGA	31 CAAGGCCACC TGCCTAGGAT CGCCCGAGT ACATTGTGT GCAGCTTTCT GCACCAGGGG GCACAGGGGC CTGGTGTCCC CAGCTGCCCA TGACCCTGA TCAATGACT TCTGCTGGTGG TGTGTCTGC TGTGTGTGC TGTGTGTG	CAGGGGGGA GCAGCCCAG GCAGCCCAG GCAGCCCAG GCAGCCCAG GCAAGGCCTG AGTGCAGTAC TGAATGCTC CAAGGCTGAAG GCAGCTGAAG GCAGCTGAAG CAGCATCTTG CAGGCTGAG CAGCATCTTG CAGGCCAAGC CAGCAAGCCAAGC	GCAAGGGACA CTTCTGGTGG CACAGGGAGA GGCTCCTCAT GTGCTGCCTT AGCGATGACC CAGGCCATCC CATGTGGCTG ATCCTGATCA GGGCAGGGGG GAGTTGCCT AGGACACCTAC ACCCGACCTC AGCCAATCCT CAGTACACTC GTCCAGCTG	120 180 240 300 360 420 540 600 720 780 840 900 960
50 55 60	Nucleic According sequences of the control of the c	Id Accession uence: 99-8: 11 GGGGCTGGGC TCCTAGCTGA CGCCGGGATT CAGCAATTTC AGCCAGTGCA GTTCGCCCAG GTCCCAGGC TGCCCAG GTCCCAGGC TGCTCCAGGC TGCTCCAGGC GTCCCAGGAC TGCTGGGGG CTCCGACTTC TCCCGGAGA GACCTCTGCT GTGGACAG GTCTGGGC GTTGCCCAG TTCCCGGACA TTCCGGACA TTCCGGACA TTCTGGGCG TTCTGGGACA TTTGCGGCTT TTTGCGGCTT	#: NM_0000 333 21 TCGGACCTGC CGGCTTTTAC CTGGCAGAGG CAGGGTGTGC CAGGGTGTGC CAGGGTGTGC CTGGCCGAC CTGGTGGACA ATCAAGAATG TTCTTCTTCG GTGTGCACGA CCACGAGGCC CCACTGCCGA CCACTGCCGAC GCCAGTGCCC CCACTGCCGA CGGGGTCCCCACGCCGACGGCCC CCACTGCCGAC CGGGGTCCCCACGCCGCCCACTGCCCA	31 CAAGGCCACC CGCCCGAGT ACATTGTGT GCAGCTTTCT GCTTTGCCAC GCTCTGGGGG GCACAGGGGC CTGGTGTCCC CAGCTGCCCA CTGACCCTGA TCAATGACTT TGGTGTCG TGGTGTCGC TGGTGTCGC TGGTGCGCA GGTGACCTGA GGTGACCTGA GGTGACCTGA GGTGACCTGA GGTGACCGGA GGCACTGAC	GCAGGGGGA GCAGGGGGGA GCCAGCCCAG GCTACTGGAT CGAAGGCTT TGATGTATCTC CCAAGGTCTGC CAAGGTCTGC CAAGGTCTGC CAAGGTCTGAAG GGAGCTGAAG CAGCATCTTG CGTGCCTGTG TGAGCCAAGC CTACAAGGTC GGAGGTGAAC	GCAAGGGACA CTTCTGGTGG CACAGGGAGA GGCTCCTCAT GTGCTGCCTT AGCGATGACC CACGGCCATCC CATGTGGCTG ATCCTGATCA GGGCAGGGGG CGAGTTGCCT AGGACACTAC ACCGACCTC AGCAATCCT CAGTACACTC GTCCCAGCTG GTCCCAGCTG	120 180 240 300 360 420 540 600 720 780 840 900 960 1020
50 55 60	Nucleic According sequences of the control of the c	Id Accession uence: 99-8: II GGCGCTGGGC TCCTAGCTGA CGCCGGGATTC CAGCAATTTC AGCAGTGCA GTTCGGCCTG GTCCCAGGG CTCCCAGGG GTCCCAGGG GTCCCAGGG GTCCCAGGG GTCCCAGGG GTCCCAGGG GTCCGACTTC TTCCCGGAGA AACCTCTGCT GTGGACAG GTGGACAGC GTTGGACAGC GCTGGACAGC GCTGGGACAG	#: NM_0000 21 TGGGACCTGC CGGCTTTTAC CTGGCAGAGG TACGCCGCTG CGCAGGTCC CAGGGTGTGC CATGCCGAC ATCAGAACACTC CTGGCCGAC CTGGTGACA ATCAAGAATG TTCTTCTTCG GTGTGACAC CCACTGCGA CCACTGCCGA CGCAGTGGCC CACTGCGAC ATCGGGGAGG CGCAGTGCCC ATCGGGGAGG	31 CAAGGCCACC TGCCTAGGAT CGCCCGAGT ACATTGTTT GCTTTGCCAC GCTCTGGGGG GCACAGGGGC CTGGTGTCCC CAGCTGCCA TCAATGACTT CTGCTGGTGG TCGTGGTGCT CTGTGACTGC GTGAGCGCA GCCACTGAC CTGTGACCGA	CAGGGGGGA GCAGGCCAG GCAAGGCCAG CTTACTGGAT CGAAGGGCTA TGATGTGATC TGCAATTCTC CAAGGTCTGC AAGGCTGAAG GGAGCTGAAG GGAGCTGAAG CAGCATCTTG CGTGCCTGTG TGAGCCAAGC CTACAAGGTC CGAGTACCAA GACAGCTCAG	GCAAGGGACA CTTCTGGTGG CACAGGGAGA GGCTCCTCAT GTGCTGCCTT AGCGATGACC CATGTGGCTG ATCCTGATCA ATCCTGATCA AGCGATGCCT AGGACACTAC AGCGACTTC AGCAATCCT AGCAATCCT CAGTACACTC CAGTACACTC CAGTACACTC AGCCAATCCT CAGTACACTC GTCCCAGCTC AGCAATCCT CAGTACACTC AGTGACTGTAA ACCACTGCCC	120 180 240 300 420 480 540 600 660 720 780 840 900 900 1020 1080
50 55 60	Nucleic According sequences of the control of the c	Id Accession lence: 99-8: II GGCGCTGGGC TCCTAGCTGA CGCGGGATC CACGGCCTT AGCAATTCA AGCAGTCCA GTTCGGCCG GTCCCAGG GTCCCAGGAC TCCTGTGGGC TCCCAGGAC TCCCGGACT TCCCGGACT TCCCGGACA GACCTCTGC GGCTGGACAG GCTGGGACAG GCTGGGACAG GCTGGGACAG GCTGGGACAG GGTGGGACAG GGGCAACAGC GGAACTGACC	#: NM_0000 21 TCGGACCTGC CGGCTTTTAC CTGGCAGAGG TACGCCGCTG CGGAGGTCC CAGGGTGTGC CAGGGTGTGC CTGGCCGAC CTGGCCGAC CTGGTGACACACTC CTGGCCGAC CTGGTGACAC CTGGTGACA ATCAAGAATT CCAGAGACC CCACTGCCGA CCACTGCCGA CGCATGCCGA CGAGGCC CCACTGCCGA ATCAGGATA ATCAGGATA ATCAGGATA ATCAGGATA ATCAGGATA ATCAGGATA ATCAGGATA ATCAGGATA ATCAGGATA	31 CAAGGCCACC TGCCTAGGAT CGCCCCGAGT ACATTGTGT GCAGCTTTCT GCACCTGAG GCACAGGGGC CTGGTGTCCC CAGCTGCCCA TCACCCTGA TCATGACCTGC TGGTGCTGC CTGTGACTGG GTGACCTGG GTGACCTGA CTGTGACCTGC CTGTGACCTGC CTGTGACCTGC CTGTGACCTGC CTGTGACCTGC CTGTGACCTGC CTGTGACCGCC CTGTGACCGCC CTGTGACCGC	CAGGGGGGA GCAGGCCAG GCAGCCCAG GCAGCCCAG GCAGCCCAG GCAATCCT CAAGGCTGAAG GCAGCTGAAG GCAGCTGAAG CAGCATACT CAGGCTGAG CAGCATACT CAGGCCTGAG CAGCATACT CAAGGCCAAG CAGCATACAG CAAGACCAAG CAAGCCAAG CAAGCCAAG CAAGCCCCCCT CAAGCCCACC CAACAGCCCCCCT CAACAGCCCACC CACACCCCCCCC	GCAAAGGACA CTTCTGGTGG CACAGGAGA GGCTCCTCAT GTGCTGCCTT AGCGATGACC CAGGCCATCC CATGTGGCTG ATCCTGATTA CGGCAGGGG CGAGTTGCCT AGGCACCTAC ACCCGACCTC AGCCAATCCT CAGTACACTC GTGACTGATA ACCACTGCC GTGACTGCC GTGACTGCC GTGACTGCC GTGACTGCC GTGACTGCC GTGGCCTGCC	120 180 240 300 420 480 540 600 660 720 780 840 900 960 1020 1080 1140
50556065	Nucleic Acc Coding sequence	Id Accession uence: 99-8: 11 GGGGCTGGGC TCCTAGCTGA CGCCGGGATT CAGCAATTTC AGCCAGTGCA GTTCGCCTG CTACAAGGG CCTGCCCAG GTCCCAGGC TTCCCAGGAC TCCTGGGCTG TTCCGGACT TCCGGACT TTCCGGACA GTCTGGGCG GCTGGCCAG GCTGGCCAG GCTGGCCAG GCTGGACA GCTGGACAG GCTAGACAGC GCAACAGC GGAACTGAC AGGTGCACC	#: NM_0000 21 TGGGACCTGC CGGCTTTTAC CTGGCAGAGG TACGCCGCT GGCAGGTCC CAGGGTGTGC CTGGCCGAC CTGGTGGACA ATCAAGAATG TTCTTCTTCG GTGTGCACGA CCACGAGGCC CCACTGCCGA CCACTGCCGA CCACTGCCGA ATCAGAGACC GCCAGTGCCA ATCAGAGACC GCCACTGCCGA ATCACGAGACA ATCACGAGAGAC ATCACGAGAC ATCACCAC ATCACCAC ATCACCAC ATCACCACAC ATCACCAC ATCACCA	31 CAAGGCCACC TGCCTAGGAT CGCCCGAGT ACATTGTGT GCAGCTTTCT GCTTTGCCAC GCTCTGGGGG GCACAGGGGC CTGACCCCA TGACCCTGA TCAATGACT TCTGTGTGCCC TGGTGTCCC TGGTGTCCC TGGTGCCGA CTGACCTGA CTGACCTGA CTGTGACTGC TGGTGCGGCA TCAATGACT TGTGTGACTGC TGTGACGGCA GCCACTGAC TGACAGCCCA TCACAGCCCA TGACATGGCG TGACAGCCCA	CAAGGGGGGA GCAGGCCCAG GCAAGCCCAG GCAAGCCCAG GCAATTCTC GAAGGCTGAAG GCAACTCTC CAAGGCTGAAG GCAGCTGAAG CAGCACTCAG CAGCACTCAG CAGCACCAG CAGCCCCCAG GCAGCCCCCAG GCACCCCCCAG GCACCCCCCAG GCACCCCCAG GCACCCCCAG GCACCCCCAG GCCCCCCAG GCCCCCCCAG GCCCCCCCAG GCCCCCCCAG GCCCCCCCAG GCCCCCCAG GCCCCCCCAG GCCCCCCCAG GCCCCCCCAG GCCCCCCCAG GCCCCCCCAG GCCCCCCAG GCCCCCCCAG GCCCCCCCAG GCCCCCCCAG GCCCCCCAG GCCCCCCCAG GCCCCCCAG GCCCCCCCAG GCCCCCCCAG GCCCCCCCAG GCCCCCCCAG GCCCCCCCC	GCAAGGGACA CTTCTGGTGG CACAGGGAGA GGCTCCTCAT GTGCTGCCTT AGCGATGACC CACGGCCATCC CATGTGGCTG ATCCTGATCA GGGAGGGG CAAGTGCCT AGGGACACC AGCCACCC AGCCACCC AGCCACCC AGCCACCC AGCCACTC AGCCACTC AGCCACTC AGCCACTC GTGCCACCC GTGCCACCC GTGCCCGC GTGCCTGC GTGCCTGC GTGCCCC GTGCCCCC GTGCCCCC	120 180 240 300 420 480 540 660 720 780 840 900 960 1020 1080 1140
50 55 60	Nucleic According sequences of the control of the c	Id Accession uence: 99-8: 11 Geocottagec TCCTAGCTGA CGCCGGATTC CAGCAATTTC AGCCAGTGCA GTTCGGCCTG GTCCCAGGG TCCCAGGGC TGCTCCAGGG GTCCCAGGG GTCCCAGGG GTCCCAGGGC TGCTGGGGGG CTGCGCCAG GTCCGACTC TCCGGAGA AGCCTTGCT GTGGACAGC GCTGGCCAG GCTGGCCAG GGTGGACAGC GGTGGACAGC GGTGGACAGC GGTGCACACAGC GAGTTGCCAC GGAGTGCCAC GGAGTGCCAC GGAGTGCCAC GGAGCTGGGC GGAGTGCCAC GGAGCTGGGC GGGCCTGGGC GGGCCTGGGC GGAGCTGGGC GGGCCTGGGC GGCCTGGCC GGGCCTGGC GGGCCC GGGCCC GGGCCC GGGCCC GGGCCC GGGCCC GGCCC GCCC GGCCC GCCC GGCCC GCCC GCC GCCC GCCC GCCC GCCC GCCC GCCC GCCC GCCC GCC GCCC GCC GC	# : NM_0000 333 21 TGGGACCTGC CGGCTTTTAC CTGGCAGAGG TACGCCGCTG CGGAGGTCC CAGGGTGTGC CAGGGTGTGC CTGGCCGAC ATCAAGAATG TTCTTCTTCG GTGTGACAC CCACTGCCGA CCACTGCCGA CCACTGCCGA CCACTGCCGA ATCAAGAATA GGCTACCGTA CCGGGAGGC CCACTGCCGA ATCAGGAGC CCACTGCCGA CCCACTGCCGA CCCACGCGCAGC CCCACTGCCGA CCCACGCGCAGC CCCACGCGCAGC CCCCGCGCAGGCC CCCCGCGCAGGCC CCCCGCGCAGGCC CCCCGCGCAGGCC CCCGCGCAGGCC CCCGCGCAGGCC CCCGCGCAGGCC CCCGCGCAGGCC CCCGCGCAGGCC CCCGCGCAGGCC CCCGCGCAGGCC CCCGCGCAGCC CCCCGCGCACC CCCCGCGCAGCC CCCCGCGCACC CCCCCGCGCACC CCCCCGCCC CCCCCCCC	31 CAAGGCCACC TGCCTAGGAT CGCCCGAGT ACATTGTTT GCTTTGCCAC GCTCTGGGGG GCACAGGGC CTGGTGTCCC CAGCTGCCA TCAATGACTT CTGCTGGTGG TCGAGGGC TGGTGCTGC TGGTGCTGC TGGTGCTGC TGGTGCTGC TGGTGCTGC TGGTGCTGC TGTGACTGC TGTGACTGC TGTGACTGC TGTGACTGC TGTGACTGC TGTGACTGC TGTGACTGAC TGTGACTGC TTCAGTGTT	CAGGGGGGA GCAGGCCAG GCAAGCCCAG CTTACTGGAT CGAAGGCTTA TGATGTGAT TGATGTAT TGATGTAT TGATGTAT TGATGTAT TGATGTAT TGATGTAT TGATGTAT TGAGGTCTGC AAGGCTGAAG CGAGCTGAAG CGAGCATCTTG TGAGCCAAGC CTACAAGGTC GGAGGTGAAC GGAGGTGAAC GAAGCTCCTG GGAGCTCCTG GGAGCTCCTG GGTCCTCTG GGTCCTCAGT GCTCCTGAC	GCAAGGGACA CTTCTGGTGG CACAGGGAGA GGCTCCTCAT GTGCTGCCTT AGCGATGACC CACGGCCATCC CATGTGGCTG ATCCTGATCA GGGACACTAC AGGACACTAC AGGACACTAC AGCCAACCTC AGGACACTAC GTCCCAGCTG GTGACTGACT ACCAGCTG GTGACTGCC GTGGCCTGC GTGGCCTGC GTGGCCTGC GTGGCCTGC GTGGCCTGC GTGGCCCC TTGGAGCCTG	120 180 240 300 360 420 660 720 840 900 900 900 1020 1140 1200 1260
50556065	Nucleic According sequences of the control of the c	Id Accession lence: 99-8: II GGCGCTGGGC TCCTAGCTGA CGCGGGATC CACGGCCTT CAGCAATTTC AGCAGTGCA GTTCGGCTG CTACAAGGG CTCCCAGGAC TCCTGCCCAG GTCCCAGGAC TCCCAGGAC TCCCGGACTTC TCCCGGAGA GACCTCTGC GGTGGGCACACAGC GCTGGGACAG CGCAACAGC GCAACAGC AGGTGCCAC AGGTGCCAC AGGTGCCACT GGAGCTGGG TGAGGTGCC	#: NM_0000 21 TCGGACCTGC CGGCTTTTAC CTGGCAGAGG TACGCCGCTG CGCAGGTGTGC CAGGGTGTGC CAGGGTGTGC CAGGGTGTGC CTGGCCGAC ATCAACACTC GTGTGACAC ATCATCACT GTGTGCACA ATCACAGAGT CCACGAGGC CCACTGCCGA CCACGAGGC CCACTGCCGA ATCACGAGGC CCACTGCCGA CGCAGTGCC CTGCCGAC CTGCCGAC CGCAGTGCC CTGCCGAC CGCAGTACCCGT CCCACTGCCGAC CGCAGTACCCGT CCCACTGCCGAC CCCACTGCCGAC CCCACTGCCGAC CCCACTGCCGAC CCCACTGCCGAC CCCACTGCCGAC CCCACTGCCGAC CCCACTGCCGAC CCCACTGCCCAC CCCACTGCCAC CCCCACTGCCCAC CCCACTGCCCAC CCCACTGCCCAC CCCACTGCCCAC CCCACTGCCAC CCCCACTGCCCAC CCCACTGCCCAC CCCACTGCCCAC CCCCACTGCCCAC CCCACTGCCCAC CCCACTGCCCAC CCCACTGCCCAC CCCACTGCCCAC CCCCACTGCCCAC CCCACTGCCCAC CCCACTGCCCAC CCCACTGCCCAC CCCACTGCCCAC CCCACTGCCCAC CCCACTGCCCCCAC CCCACTGCCCAC CCCACTGCCCCAC CCCACTGCCCAC CCCACTGCCCCAC CCCCACTGCCCAC CCCACTGCCCCAC CCCACTGCCCCAC CCCACTGCCCCAC CCCCACTCCCCAC CCCCCACTCCCCCCCCCC	31 CAAGGCCACC TGCCTAGGAT CGCCCCGAGT ACATTGTGT GCACCTTTTGCCAC GCTCTGGGGG GCACAGGGGC CTGATGTCCC CAGCTGCCA TGACCTTGA TCAATGACTT CTGCTGGTGG GTGACCTGA CTGAGCGGC CTGAGCGGC CTGAGCGGC CTGAGCGGC CTGAGCGGC CTGAGCGGC CTATGAGCGG CTATGAGCGG TTATTTGGCCG GTTCAGTGT TATTTGGCCG	CAGGGGGGA GCAGCCCAG GCAGCCCCCG GCAGCCCCCCG GCCCCCCCC	GCAAAGGACA CTTCTGGTGG CACAGGAGA GGCTCCTCAT GTGCTGCCTT AGCGATGACC CAGGCCATCC CATGTGGCTG ATCCTGATCA GGGCAGGGG GAGTTGCCT AGGCAACTC ACCCGACCTC AGCCAATCCT CAGTACACTC GTCCAGCTG GTGACTGAC TCGGCCTGGC GTGGCCTGGC GTGGCCTGC GTGGGCCTGC GTGGGCCTGC CCCGCCACTT	120 180 240 300 360 420 600 660 720 840 900 900 900 1020 1140 1200 1260 1320
50556065	Nucleic According sequences of the control of the c	Id Accession uence: 99-85 11 GGGGCTGGGC TCCTAGCTGA CGCCGGGATT CAGCAGTTTC AGCCAGTGCA GTTCGCAGTGCA GTCCAGGGCT TCTGCCCAG GTCCCAGGAC TCCCGAGAC TCCCGACATTC TCCCGAGAC GTCCCGACAC GTGCCACACGC GGAACTGCA CGCAACAGC GGAACTGACC TGGGACAC TGTGGGACA TGTGGGACA TGTGGGCTG TGTGGGACA TGTGGGACA TGTGGGACA TGTGGGACA TGTGGACAC TGTGGACAC TGAGCTGAC TGAGGTGAC TGAGGTGAC TGCACTGAC TTGCACTGAC TTGCACTGAC TGAGGTGAC TGCACTGAC TTGCACTGAC TTGCACTGAC TCGCACTGAC	#: NM_0000 333 21 TGGGACCTGC CGGCTTTTAC CTGGCAGAGG CAGGGTGTGC CAGGGTGTGC CAGGGTGTGC CTGGCCGAC CTGGTGGACACTC CTGGCCGAC CTGGTGGAC ATCAAGAATG TTCTTCTTCG CCAGTGCCC CACTGCCGA CCACTGCCGA ATCAGAATA TCCAGAATA CGCACTGCCGA CGCAGTGCCC CTGGTGGAC ATCAGGAGAC CCACTGCCGA CGCAGTGCCC CTGCGCGAGAC CCCCCCCTCCTGTGCCCGA CGGGAGAC CCCGGAGACC CCACTGCCGA CGGGAGAC CCCGGACACC CCTTCTGTTG	31 CAAGGCCACC CGCCCGAGT ACATTGTGTT GCAGCTTTCT GCTTTGGCAC GCTCTGGGGG GCACAGGGGC CTGGTGCCCA CTGACCTGA TCAATGACT TCTGTGGTGG TCTGGTGG TCTGGTGGG TCTGGTGGG TCTGGTGGG TGTGACTGG TGTGACTGG TGTGACTGG TGTGACTGG TGTGACTGG TTGACTGG TTGACTGG TTGACTGG TTGACTGG TTGACTGG TTGACTGG TTGACTGG TTGACTGG TTGACTGG TTATTGGCCG ACAGCCCA TGACATGCC TTATTTGGCCG ACAGACCCT TATTTGGCCG ACAGACCCT	CAGGGGGGA GCAGGGGGGA GCAGCCCAG GCAGCCCAG GCAGCCCAG GCAGCCCAG CTTACTGGAT CGAAGGCTG AGGCTGAAG CAAGCTGAAG CAAGCTGAAG CAGCACTTAC CAGGCTGAAG CAGCACTTAC CAGCACTCAG CAACCCCAG CAACCCCAG CAGCTCAG GCAGCTCAG GCAGCTCAG CAGCTCCAG CAGCTCCAG CAGCTCCAG CAGCTCCAG CAGCTCCAG CAGCTCCAG GCTCCCTCAGT GCTCCCTCAGT GCTCCCTGCGTGAC CAGTTCCGGG CCCCCGGTCCCCGGT	GCAAGGGACA CTTCTGGTGG CACAGGGAGA GGCTCCTCAT AGCGATGACC CGCGCCATCC CATGTGGCTG ATCCTGATCA ACCCGACCTC AGGCAATCC CAGGCAGGGG CGAGTTGCCT AGGCAATCCT AGGCAATCCT AGCCAACCTC AGCCAATCCT CAGTACACTC GTCCAGCTG GTGACTGGACCTG CTGACCTGGCCACTT ACCCGGCCCACTT ACCCGCCACTT ACCCGCCACTT ATCCTGGGCCCA	120 180 240 420 420 420 540 600 540 720 780 960 1020 1140 1260 1320 1380
5055606570	Nucleic According sequences of the control of the c	Id Accession uence: 99-8: 11 Geocottagec Tectagetga Tectagetgat Caccaggatt Caccaggatt Caccaggatt Caccagtgat Tectagetga Geocottagec Teccactagec Tecc	# : NM_0000 333 21 TGGGACCTGC CGGCTTTTAC CTGGCAGAGG TACGCCGCTG CGGAGGTCC CAGGGTGTGC CAGGGTGTGC CTGGCCGAC ATCAAGAATG TTCTTCTTCG GTGTGACAC CCACTGCCGA CCACTGCCGA CCACTGCCGA ATCAAGAATA GCTACCGGGAGG CCACTGCCGA ATCAGGAGCC CCACTGCCGA CCCACTGCCGA CCCACTGCCGC CTTCCGGGAGG CCTGGGCAGG CCTGCGGAATA GGCTACCGTG CCTGGGCAGG GTGAGCACCC GCTTCTGTTG TGGAACTTGG	31 CAAGGCCACC TGCCTAGGAT CGCCCGAGT ACATTGTTT GCTTTGCCAC GCTCTGGGGG GCACAGGGCC CAGCTGCCAA TCAATGACTT CTGCTGGTGG TGGTGTGTCC CTGTGTGTGC CTGTGACTGC GTCAATGACTG GTCAACGCCA GCCACTGAC GCCACAGCCA TGACTGAC TTATTTGGCCG TTCAGTGTT TATTTGGCCG AGCAGACCCT TGCCTGAGGCC TGCCTGAGGCC TGCCTGAGGCC TGCCTGAGGCC TGCCTGAGGCC TGCCTGAGGCC TGCCTGAGGCC	CACAGGGGGGA GCAGGGGGGGA GCAGGCCCAG GCAGCCCAG GCAGCAGT CGAAGGCTG CGAGGTCT CCAAGGTCTC CCAAGGTCTC CCAAGGTCTGC CAAGGCTGAAG GCAGCTGAAG CAGCATCTTG CGAGCTGAAG CAGCATCTTG CGAGGTGAAG GCACATCTG GCAGGTGAAC GCAGCTCCAG GCAGCTCCAG GCTCCAGT GCTGCGTGAC CAGTTGCGGGCCCCGTGAC CAGTTGTGGGGCCCCGGTGAC CAGTTGTGGGGCCCCGGTGAC CAGTTGTGGGGCCCCGGTCCCCGGGCCCCGGTAC CCCTGGCTAC	GCAAGGGACA CTTCTGGTGG CACAGGGAGA GGCTCCTCAT GTGCTGCCTT AGCGATGACC CACGGCCATCC CATGTGGCTG ATCCTGATCA GGGACGCGG CGAGTTGCCT AGGACACTCC AGGACACTCC AGCAATCCT AGCAATCCT CAGTACACTC GTCCCAGCTG GTGACTGCC GTGGCTGGAC TTGGACCTG CCGCCACTT ATCCTGGGCC CGGTTGGAAT	120 180 240 300 360 420 600 660 720 840 900 900 900 1020 1140 1200 1260 1320
5055606570	Nucleic According sequences of the control of the c	Id Accession lence: 99-8: II GGCGCTGGGC TCCTAGCTGA CGCGGGATTC CACGGGCTT AGCAATTTC AGCAGTGCA GTCCCAGGAC GTCCCAGGAC TCTCCAGGAC TCTCCAGGAC TCTCGGACA TCTCGGACA GTCCAGACAGC GCTGGACAGC GCTGGACAG GCTGGGACAG CGCAACAGC GGAACTGAC CGCACTGC CCCCTTTC CGACTGGC CCTCTTTC CGACTGC CCTCTTTC CGACTGGC CCTCTTTC CGACTGGC CCTCTTTC CGACTGGC CCTCTTTC CGACTGGC CCTCTTTC CGACTGGCTTTC	#: NM_0000 21 TCGGACCTGC CGGCTTTTAC CTGGCAGAGG TACGCCGCTG CGCAGGTGCC CAGGGTGTGC GATGCACTC CTGGCCGAC CTGGTGGACA ATCAAGAATG TTCTTCTTCG GTGTGACAC CCACGAGAC CCACGAGAC CCACGAGAC CCACTGCCGA CCACTGCCGAC CCTGCACCCC CCTCTCTTTG CGACCACCGC	31 CAAGGCCACC TGCCTAGGAT CGCCCCGAGT ACATTGTGT GCTTTGCCAC GCTCTGGGGG GCACAGGGGC CTGGTGTCCC CAGCTGCCA TGACCTGA TCAATGACTT CTGCTGGTGG GTGACCTGA CCTGACCTGA	CAGGGGGGA GCAGGGGGGA GCGAGCCCAG GCGAGCCCAG GCGAGCCCAG AGTGCAGTAC CAAGGCTGC AAGGCTGAAG CAGACTGAAG CAGACTGAAG CAGACTGAAG CAAGCCAGG CTACAAGGC GCAGGTGAAC CAAGTACCAA GCAGCTCCTG GCTCCTCG GCTCCTCG GCTCCTCG GCTCCTGG CAGCTCAGT CCAGGTCCCGGTC CCGTGGCTAC ACTGCCGGTC CCGTGGCTAC ACTGCCCTCT ACTGCCCTCC	GCAAGGGACA CTTCTGGTGG CACAGGGAGA GGCTCCTCAT AGCGATGACC CAGGGCATCC CATGTGGCTG ATCCTGATCA AGCGAGGGG CGAGTTGCCT AGGACACTCC AGCCAATCCC AGCCAATCCT CAGTACACTC GTGACTGACCT GTGACTGACCT GTGACTGCC GTGACTGCC GTGACTGCC GTGGCCTGC GTGGCCTGC GGTGGCCCACTT ATCCTGGGCC CCGGTGGGAT GATGTGACCC GGTGGAAT GATGTGACCC	120 180 240 300 360 420 600 600 720 780 840 900 1020 1080 1140 1260 1320 1380 1440
50556065	Nucleic According sequences of the control of the c	Id Accession uence: 99-8: 11 GGGGCTGGGC TCCTAGCTGA GGCGGATTC CAGCAGTTCC AGCAGTTTC AGCCAGTGCA GTTCGGCCTG GTCCCAGG CTGGGCACT GTGGACAGC GGTGGACAGC GGTGGCCAACAGC GGTGCCACACAGC GGAGTGCCC TCGCACTGC TCGCACTGC TCGCACTGC TCGCACTGC TCGCACTGC TCGCACTGC TCGCACTGC TCGCACTGC TCGCACTGC GAGTGCCC GACTGGCTC GGACTGGC GAGTGGCCT GGACTGGC GAGTGGCCT CGACTGGC CCACAGGC GGATGGCCT CGCACTGC CCTCCTTTCC GACTGGCCT CGAGTGGCCT CGAGTGGCCT CGAGTGGCC CGAGTGGCCT CGAGTGGCC CGAGTGGCCT CGAGTGGCC CGAGTGGCCT CGAGTGGCC CGAGTGCC CGAGTGGCC CGAGTGGCC CGAGTGGCC CGAGTGGCC CGAGTGGCC CGAGTGGCC CGACTGC CGAGTGGCC CGAGTGCC CGAGTGCC CGAGTGCC CGAGTGCC CGAGTGCC CGAGTGCC CGAGTGCC CGAGTGCC CGAGTGCC CGCACTGC CGAGTGCC CGCACTGC CGCC CGC	# : NM_0000 333 21 TGGGACCTGC CGGCTTTTAC CTGGCAGAGG TACGCCGCG GATGCACTTG GGCAGGTCC CAGGGTGTGC CTGGCCGAC CTGGTGGACA ATCAAGAATG TTCTTCTTCG GTGTGCACGA CCACTGCGA ACCACTGCGA ACCACTGCGA ATCAGAGAC GCCAGTGCCC ATCGGGAGG CCACTGCGA ATCAGAGAC GCCAGTGCCC ATCGGGAGG ATCAGGAGC CCACTGCGA ATCAGGATC CCTGGGCAGG TGGACCACGC GCTTCTGTTG TGGAACTTGG GAGCCACGC CAGCCGGCA ACCCCTGCAA	31 CAAGGCCACC TGCCTAGGAT CGCCCGAGT ACATTGTTT GCTTTGCCAC GCTCTGGGGG GCACAGGGGC CTGGTGTCCC CAGCTGCCA TCAATGACTT CTGCTGGTGG TCAATGACTT CTGCTGGTGG CCACAGCGCA GCCACTGACGGC GTCAGGCGC GTTCAGTGGC GTTCAGTGGC GTTCAGTGGC GTTCAGTGT TATTTGGCCG AGCAGACCCT AGCAGACCCT AGCAGACCCT AGCAGACCCT CCCTGAGGC CCACAGCCC CCCTGAGGC CCCCTGAGGC CCACAGCCC CCCTGAGGC CCCCTGAGGC CCCCTGAGGC CCCCTGAGGC CCCCTGAGTCC CCCTGGTTCC CCGTGGTTCC	CACTGCCTC CACTGCACT CACTGCAG CACCCAG CTTACTGGAT CGAAGGCTG CGAGGCTGAG CAGGCTGAAG CAGCTGAAG CAGCTGAAG CAGCTGAAG CAGCATCTTG CGTGCCTGTG TGAGCCAAGC CTACAAGGTC CGAGGTGAAC CAGCTCCTG GACGTCCTG GATCCTCAGT GCTGCGTGAC CAGTTGCGGTGAC CAGTTGCGGTGAC CAGTTGCGGTGAC CAGTTGCGGTGAC CAGTTGCGGTGAC CAGTTGCGCTCC CCTCACACTC CCTCACACTC	GCAAGGGACA CTTCTGGTGG CACAGGGAGA GGCTCCTCAT GTGCTGCCTT AGCGATGACC CGCGCCATCC CATGTGGCTG ACCAGGGAGG GGAGAGGGG CGAGTTGCCT AGCGACCTC AGCCACCTC AGCCACCTC AGCCACCTC AGCCACTC CTCCAGCTG GTGACTGTGA ACCAGTGCCG GTGGCCTGC CTGGCCTGC CTGGCCTGC CCGCCACTT ACCTGGGCCCA TTGGAGCCTG CCGCCACTT ATCCTGGGCCC CGGTTGGAAT GATGTGACCC CGGTTGGAAT GATGTGACCC CACCTCTG	120 180 240 300 360 420 540 600 720 780 840 960 1020 1080 1140 1260 1320 1380 1450
5055606570	Nucleic According sequences of the control of the c	Id Accession lence: 99-8: II GGCGCTGGGC TCCTAGCTGA CGCCGGATTC CACGGCTT CACGATTCA AGCAGTGCA GTCCCAGGAC CTCCCAGGAC TCCCAGGAC TCCCAGGAC TCCCAGGAC TCCCAGGAC TCCCAGGAC TCCCAGACATC TCCCGGACA GCTCGCCAC GCTGCGCCAC GCTGCGCCAC GCTGCGCCC TGTGGACAGC GCAACAGC GGAACTGCC TGAGGTGCCACT TGAGGTGCCACT TGAGGTGCCC TGAGGTGCCC TGAGGTGCCC TGAGGTGCC TGAGGTGCC TCCCTTTCC GACTGGCTTG CCACTGCC CACTGGCTTC CCACTGCC CACTGGCTTC CACGGCTTGC CACTGGCTTC CACGGCTTGC CACGGCTTGC CACAGGCCTGC CACAGGCCTGC CACAGGCCTGC CACAGGCCTGC CAACAGCCTGC CAACAGCCTGC CAACAGCCTGC CAACAGCCTGC CAACAGCCTGC CAACAGCCTGC CAACAGCCTGCC CAACAGCCTGC AACAGACCTG	#: NM_0000 333 21 TGGGACCTGC CGGCTTTTAC CTGGCAGAGG TACGCCGCTG CAGGGTGTC CAGGGTGTC CAGGGTGTC CTGGCCGAC CTGGCCGAC CTGGTGACAC CTGGTGACAG CCACGAGACC CCACTGCCGA CCACTGCCGA CCACTGCCGA CCACTGCCGA CCACTGCCGA ATCAGAATA GCTACCGTTC CTGGCAGACC CCACTGCCGA CTGGGACC CCACTGCCGA ATCAGAGAC GCTACCGTC CCTGCGAAC CCACTGCCGC CATCCCGC CTGCGACCC CCTGCCGA CTGCCGC CAGCCGCC CAGCCGCC CAGCCGCC CAGCCGCC CAGCCGCCAACCACCCC	31 CAAGGCCACC TGCCTAGGAT CGCCCCGAGT ACATTGTGT GCTTTGCCAC GCTCTGGGGG GCACAGGGGC CTGGTGTCCC CAGCTGCCA TGACCTGA TCAATGACTT CTGTGGGGG GTGACCTGA CTGACCTGA CTGACCTGA GCACATGAC TTGACTGGC GTGACCAC GGCCACTGAC CTGAGCGCCA GGCCACTGAC CTGAGCGCCA TGACATGGCG GTCAGTGGT TATTTGGCCG AGCAGACCCT TGCCTGAGGC TGAGTGGT TGCTTGAGGCG CCATGACC AGCTGATCC AGCTGGTTCC AGCTGCCCGG	CAGGGGGGA GCAGGGGGGA GCGAGCCCAG GCAGCCCAG GCAGCCCAG GCAGCCCAGCCCAGCCCCCCCC	GCAAGGGACA CTTCTGGTGG CACAGGGAGA GGCTCCTCAT AGCGATGACC CAGGGCATCC CATGTGGCTG ATCCTGATCA AGCCAGCGCC CGAGTACC CGAGTACC CGAGTACC CGAGTACC CGAGTACC AGCCACTC AGCCAATCCT CAGTACACTC GTGACTGAC ACCAGCTG CTGACTGAC ACCAGCTG CTGACTGCC GTGGCCTGC CGTGGCCTGC CGTGGCCTGC CGGTGGAAT ATCCTGGGCC CGGTTGGACC CGGTTGGACC CGGTTGGACC CGGTTGGACC CGGTTGACC CGGTTGCACC CGGTTGCACC CGGTTGCACC CGGTTGCACC CGGTTGCACC CGGTTGCACC CGGTTGCCC CGGTTGCACC CGGTTGCCC CGGTGCCC CGGTTGCCC CGGTTGCC CGGTTGCC CGGTTGCC CGCC CGGTTGCC CGGTTGCC CGGTTGCC CGGTTGCC CGGTTCC CGCC CGGTTGCC CGCC	120 180 240 300 360 420 540 600 720 780 840 900 1080 1140 1260 1320 1380 1440 1500 1500 1680
5055606570	Nucleic According sequences of the control of the c	Id Accession lence: 99-8: II GGCGCTGGGC TCCTAGCTGA CGCGGGATC CACGGCCTT CAGCAATTTC AGCAGTGCA GTTCGGCCTG GTTCCCAGGAC TCCCAGGAC TCCCAGGAC TCCCAGGAC TCCCAGGAC TCCCAGGAC TCCCAGGAC TCCCAGGAC TCCCGGACTTC TCCCGGAGAC GCTGCGCCAC GGTGCCACAC GGTGCCACAC GGTGCCACAC GGAACTGAC GGAACTGAC CGCACTGC TCGCACTGC TCCACTGCC TCCACGGCC TCCACGGCC TCCACGGCC TCCACGGCC TCCACGGCC TCCACGGCC TCCCCTGTGCC TCCCTTTCC TCCCCTGTGCC TCCCTGTGCC TCCCTGTGCC TCCCTGTGCC TCCCTGTGCC TCCCTGTGCC TCCCTGTGCC TCCCTGGTGCC TCCCTGGTGCC TCCCTGTGCC TCCCTGCT TCCCTTCT TCCTTCT TCCTTCT TCCCTTCT TCCTTCT TCCTTTCT TCCTTCT TCCTTTCT TCCTTTCT TCCTTCT TCCTTTCT TCCTTTTCT TCCTTTTCT TCCTTTTCT TCCTTTTTT	#: NM_0000 21 TCGGACCTGC CGGCTTTTAC CTGGCAGAGG TACGCCGCTG CAGGGTGTGC CAGGGTGTGC CAGGGTGTGC CAGGGTGTGC CTGGCCGAC CTGGCCGAC CTGGTGAAATA TCATCATCTTCT GTGTGACAG CCACGAGACC CCACTGCCGA CCACTGCCGA CCACTGCCGA CCACTGCCGA ATCAGAATA GCTACCGTG CCACTGCCGA CTGGGCAG CCTGCGAC CCTGGCAGA ACCCCTGCAA ACCCCTGCAA ACCCCTGCAC ACCCCCTGCAC ACCCCTGCAC ACCCCCTGCAC ACCCCCTGCAC ACCCCCTGCAC ACCCCCTGCAC ACCCCAC ACCCCAC ACCCCCTGCAC ACCCCTGCAC ACCCCCTGCAC ACCCCAC ACCCCTGCAC ACCCCCTGCAC ACCCCTGCAC ACCCCCTGCAC ACCCCTGCAC ACCCCTCCAC ACCCCTCCCCCCCCCC	31 CAAGGCCACC TGCCTAGGAT CGCCCCGAGT ACATTGTTT GCAGCTTTTGCCAC GCTCTGGGGG GCACAGGGGC CTGGTGTCCC CAGCTGCCA TGACCTGA TCAATGACTT CTGCTGGTGG GTGACCTGA CCTGACCTGA	CAGGGGGGA GCAGGGGGGA GCAGCCCAG GCAGCCCAG GCAGCCCAG GCAGCCCAG GCAGCCCAG GCAGCCCAG CAGCCCAG CAGCCCAG CAGCCCAG CAGCCCCCG GCACCCCCC GCACCCCCCCC	GCAAAGGACA CTTCTGGTGG CACAGGAGA GGCTCCTCAT GTGCTGCCTT AGCGATGACC CGCGCCATCC CATGTGGCTG ATCCTGATCA CGGCAGCGG CGAGTTGCC AGCCACTC AGCCAACTC ACCCGACCTC AGCCAACTC GTGACTGATCA ACCACTGCC GTGACTGAC GTGACTGCC GTGACTGCC GTGGCCTGC GGTGGCCTGC GGTGGGCCTGC CGGTTGGACT ATCCTGGGCC CGGTTGGACCT CAGTACCTC CGGTTGCACT CCGGCCACTT ATCCTGGGCC CGGTTGGACCT CAGTGCCC CGGTTGGACCT CAGGCGCTC CAGCTGCCC CGGTTGGACC CGGTTGCACT CAGGCGCTTC CAGGCGCTTC CAGGCGCTTC CAGGCGCTTC CAGGGGGTTC CAGGGGGTTC CAGGGGGTTG	120 180 240 300 360 420 540 600 660 720 780 900 960 1020 1080 1140 1200 1320 1380 1440 1560 1560 1680 1680
505560657075	Nucleic According sequences of the control of the c	Id Accession uence: 99-8: 11 GGGGCTGGGC TCCTAGCTGA CGCCGGGATT CAGCAATTTC AGCCAGTGCA GTTCGGCCTG GTCCCAGGG CCTACCAGGGC CTACAAGGGG CCTGCCCAG GTCCCAGGA GTCCCAGGACT TCCGGACTG GTGGACAG GCTGGCCAG GCTGGCCAG GCTGGCCAG GCTGGCCAG GCTGGCCAG GCTGGCCAG GGAGTGAC GGAGTGAC GGAGTGAC GGAGTGAC GGAGTGAC CCCCCTTTCC GACTGC GAGTGGCT GCAACAGC GGATGGCT CCCCCTTTCC GACTGC CCTCCTTTCC GACTGCC CCAGGTGCC CCACAGACCT GGAGTGGCC GACAGACCT GGATGGCCT GGACTGCC CCACTGCTCC CCACTGCTCC CCACTGCTCC CCACTGCTCC CCACTGCTCC CCACTGCTCC CCACTGCTCC CCTGGTGCC ACAGACCTC GGTGCTCC GGTGCTCC GGTGCTTCC GGTGCTCC GGTGCTCC GTGCTTCC GGTGCTCC GGTGCTTCC GGTGCTCC GGTGCTTCC GGTGCTTCC TCCACTTCC GGTGCTTCC GGTGCTTCC GGTGCTCC GGTGCTTCC TCCACTTCC GGTGCTTCC TCCCTGCTCC GGTGCTTCC TCCCTCTCC GGTGCTTCC TCCCTCTCC TCCCCTCTCC GGTGCTTCC TCCCCTCTCC TCCCCCTCTCC TCCCCCC	# : NM_0000 333 21 TGGGACCTGC CGGCTTTTAC CTGGCAGAGG CAGGGTCC CAGGGTTGCC CAGGGTTGCC CAGGGTGGC CTGGCCGAC CTGGCCGAC CTGGCCGAC CTGGCCGAC CTGGCCGAC CTGGCCGAC CCACTGCCGA CCACTGCCGA ATCAAGAATG TCTTCTTCG GTGACCCGA CCACTGCCGA ATCAGAGAC CCACTGCCGA ATCAGAGAC CCACTGCCGC ATCGGGCAGG CCTCGGCAGAC CCCCCCCCCC	31 CAAGGCCACC CGCCCGAGT CGCCCGAGT ACATTGTGT GCAGCTTTCT GCACTTGCCAC GCTCTGGGGG GCACAGGGGC CTGGTGCCCA CTGACCTGA CTGACCTGA CTGACCTGA CTGACCTGA CTGACCTGA CTGACAGGGCA CTGACAGGCCA CTGACAGGCCA CTGACAGCCCA CTGACAGCCCA CTGACAGCCCA CTGACAGCCCA CTGACAGCCCA CCACAGCCCA CCACACCCCA CCACACCCAC CCACACCCCAC CCACACCCAC CCACACCCCAC CCACACCCAC CCACACCAC	CAGGGGGGA GCAGGGGGGA GCAGCCCAG GCAGCCCAG GCAGCCCAG GCAGCCCAG GCAACTCTC CAAGGCTGAAG GCAGCTGAAG GCAGCTGAAG GCAGCTGAAG GCAGCTGAAG GCAGCTGAAG GCAGCTGAAG GCAGCTCAG GCAGCTCAG GCAGCTCAG GCAGCTCAG GCAGCTCAG GCAGCTCAG GCAGCCCGGT CCTCAGT GCTGCGTGA CCAGCTCAG GCAGCACC CCTGGACCA GCAGCAGCAC GCAGCAGCAC GCAGCACC CTTGGACCA CCTTGGACCA CCTTGGACCAC CCTTGGACCAC CCTTGGATGACC	GCAAAGGACA CTTCTGGTGG CACAGGGAGA GGCTCCTCAT AGCGATGACC CGCGCCATCC CATGTGGCTG ATCCTGATCA GGGAGGAGG GGAGTGCCT AGGACATCA ACCCGACCTC AGCCAATCC CAGCTACC AGCCAATCC AGCCAATCC AGCCAATCC AGCCAATCCT CAGTACACT ACCCGACCTC AGCCAATCCT CTCCAGCTG CTGACCTGG CGTGGCCCA TTGGAGCCTG CGGCACTT ACCTGGGCC CGGTTGGAAT CATGTGACCT CAGGTGCCC CGGTTGGAAT CATCTCTCC CAGGTGCCC CGGTTGGATC CAGGTGCCT CAGGGGTTC CAGGGGTTC CAGGGGTTC CAGGGGTTC GTTCAGGCTG GTTCAGCTG	120 180 240 300 360 420 540 660 720 780 900 900 1080 1140 1260 1320 1380 1440 1500 1620 1680 1740
5055606570	Nucleic According sequences of the control of the c	Id Accession uence: 99-8: 11 GGCGCTGGGC TCCTAGCTGA CGCCGGATTC CAGCAATTTC AGCCAGTGCA GTTCGCCTG CTACAAGGGG CTGCCCAG GTCCCAGGAC TCCGACTTC TCCGGAGAC TCCGACTTC TCCGGAGAC TGCGCCAG GCTGGCACAGC GGTGGGACAG GGTGCAACAGC GGAACTGACC AGGTGCACT TGAGTGCACT GGAGTGCACT CGCACTGC CGCACTGCC GATGGCTGC CCCCATTCC GAACTGCC GATGGCTGC CCCCGTTCC GACTGCC GACTGCC CCCCCTCTTCC CCACTGCC CCCCGTTCC CCACTGCC CCCGGTGCC CCCGGTGCC CCCTGGTGCC CCCTGGTGCC CCCTGGTGCC CCTCGTTCC CCCTGTGCC CCTCTTCC CCCTGGTGCC CCCTGTGCCC GCTGTTCC CACTGTGCCG CCTCTTCCC CACTGTGCCG CCTCTTCCC CACTGTGCCG CCTCTTCCC CACTGTGCCG CCTCTTCCT CACTGTGCCG	# : NM_0000 333 21 TGGGACCTGC CGGCTTTTAC CTGGCAGAGG CAGGGTCC CAGGGTGTGC CAGGGTGTGC CAGGGTGTGC CTGGCCGAC CTGGCCGAC CTGGCCGAC CTGGCCGAC CTGGCCGAC CTGGCCGAC CTGGCGAC CTGGCCGAC CCACTGCGA CCACTGCGA ATCAGAGATG CTCACTACTG CGCACTCCGA ATCAGGACAC CCTGGCAC CTGCGCAC CTGGCAC CTGGCAC CCACTGCGA ATCAGGAC CCACTGCGA ATCAGGAC CCACTGCGA ACCCCTGCAA CAGCCACC ACCCAGTACC ACCCAGTA	31 CAAGGCCACC TGCCTAGGAT CGCCCCGAGT ACATTGTTT GCTTTGCCAC GCTCTGGGGG GCACAGGGCC CAGCTGCCCA CTGACCTTG TCGTGTGCC TGGTGTCC TGGTGTCC TGGTGTCT TGCTGGGGG GTCAGCCTGA TGACCTGA TGACCTGA TGACCACGA GGCCACTGA TGACATGCC TGACAGCCCA TGACATGCC TGACAGCCCA AGCAGACCCT TATTTGGCCG AGCAGACCCT TGCCTGAGGC AGCAGACCCT CACAGCCCA AGCAGACCCT TGCCTGAGGC AGCAGACCCT TGCCTGAGGC AGCAGACCCT CACAGCCCA AGCAGCCCA AGCAGCCCA AGCAGCCCA CACAGCCCA AGCAGCCCA AGCAGCCCA AGCAGCCCA CACAGCCCA AGCAGCCCA AGCAGCCCA CACAGCCCA CACAGCCCA AGCAGCCCA AGCAGCCCA CACAGCCCA AGCAGCCCA CACAGCCCA CACAGCCCA CACAGCCCA AGCAGCCCA CACAGCCCA CACAGCCCA CACAGCCCA AGCAGCCCA CACAGCCCA CACAGCCA CACAGCCCA CACAGCCCA CACAGCCA CAC	CAGGGGGGA GCAGCCCAG GCAGCCCAG GCAGCCCAG GCAGCCCAG GCAGCCCAG AGTGCAGTAC CAAGGCTGC AGGGCTGC AGGCTGAAG GCAGCTGAAG GCAGCTGAAG CTACAAGCTCTG CAAGGTCTGC CAAGGCTAAG CCAAGCCCAGC CACCCGTGAAG CCACCGGGTGAC CAGCTCCTG GCTCCTAGCCCAGC CCGTGGCTGC CCGTGGCTGC CCGTGGCTCC CCTCACACTC CCACCGGGTAC CCACTGGACCC CCTTGGACCAC CCTTGGATGCC CCTTGGATGCC CCTTGGATGAC CCGTGGCTAC CCTTGGATGAC CCTTGGATGAC CCTTGGATGAC CCTTGGATGAC CCGTGAGGGGC CCCTTGGATGAC CCGTGAGGGG	GCAAGGGACA CTTCTGGTGG CACAGGGAGA GGCTCCTCAT GGGCATGCC CGCGCCATCC CAGGCATCA GGCAGGGG CGAGTTGCCT AGGCACACCT AGCCACTCC AGCCACTCC AGCCACTCC AGCCACTCC AGCCACTCC AGCCACTCC AGCCACTCC CTGACTGGCCT ATCCTGACTG ACCACTCC CTGCCAGCTG CTGACTGGCC CGGTGGCCCACTT ATCCTGGGC CGGTTGGACT CACGCCC CGGTTGGACT CAGGCGTG CCGGCCACTT ATCCTGGGCC CGGTTGGCCTG CAGGGGGTTG CAGGGGGTTG CGGGGGTTG CGGGGGTTG CGGGCTGGCCTG CGGGGGTTG CGGGCTGGCCTG CGGGGGTTG CGGGCTGGCCTG CGGGGGTTG CGGGCTG CGGGTGCCTG CGGGGGTTG CTCAGGCTG ATCGCGCTG AGGGGGTTG ATCCAGCTG AGTGCCCAGTG AGCGCCAGTG AGTGCCCAGTG AGCGCCCAGTG AGTGCCCAGTG AGTGCCCAGTG AGTGCCCAGTG AGCGCCCAGTG AGCGCCCAGTG AGCGCCCAGTG AGCGCCCAGTG AGCGCCCAGTG AGTGCCCAGTG AGCGCCCAGTG AGCGCCCAGTG AGCGCCCAGTG AGCGCCCAGTG AGCCCAGTG AGCCCCAGTG AGCCCCCCAGTG AGCCCCAGTG AGCCCCAGTG AGCCCCCCAGTG AGCCCCAGTG AGCCCCCAGTG AGCCCCAGTG AGCCCCAGTG AGCCCCAGTG AGCCCCAGTG AGCCCCAGTG AGCCCCAGTG AGCCCCAGTG AGCCCCCCAGTG AGCCCCAGTG AGCCCAGTG AGCCCCAGTG AGCCCCAGTG AGCCCAGTG	120 180 240 300 360 420 540 660 720 780 840 900 900 1080 1140 1260 1380 1560 1560 1680 1740 1680 1740 1860
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505560657075	Nucleic According sequences of the control of the c	Id Accession lence: 99-8: 11 GGGGCTGGGC TCCTAGCTGA CGCCGGATTC CAGCAATTTC AGCCAGTGCA GTTCGGCCTG GTCCCAGGG TCCTAGCAGGG TCCCAGGG GTCCCAGGG TCCCAGGG GTCCCAGG GTCCCAGG GTCCCAGG GTCCCAGG GTCCAGGACT TCCGGAGA TGCGGCGG GGGGCG GGGGCGG GGGGCGG GGGGCGG GGGGCGG TGAGGTGCC TGGGCACT CGCACTGC TGGGCACT TGCCACTGC TGGGCACT TGCCACTGC TGGGCCC TGGGCCC TGGGCCC TGGGCCC TGGGCCC TGGGCCC TCGCCTGTTCC GACTGGC CCCGGTGCC CCACTGTCC CCTGTTCC GACTGGCC CCACTGGCC CCACTGGCC CCACTGGCC CCACTGGCC CCACTGGCC CCACTGGCCC GGTGCTCC CCCCGGGGAG CCGCGGGAG GAGCCACAGGCC CCCCGGGGAG GAGCCACAGGCC CCCCGGGGAG GAGCCACAGGC CCCCGGGAGT GAGCCACAGGC GAGCCACAGGC CCCCGGGAGT GAGCCACAGGC CCCGGGAGT GAGCCACAGGC CCCCGGGAGT GAGCCACAGGC CCCCGGGAGT GAGCCACAGGC CCCCGGGAGT GAGCCACAGGC CCCCGGGAGT GAGCACAGGC	# : NM_0000 333 21 TGGGACCTGC CGGCTTTTAC CTGGCAGAGG TACGCCGCTG CAGGGTGTGC CAGGGTGTGC CTGGCCGAC ATCAGACATT CTGGCAGAC CTGGCCGAC ATCAGAGAC CCAGTGCCC ATCAGAGAC CCACTGCGA ATCAGAGAC CCACTGCGA ATCAGAGAC CCACTGCGA ATCAGAGAC CCACTGCGG ATCAGGGC CCATTCTGTT GGCAGAC CTGGCAGAC CCACTGCGA ATCAGAGAC CCCGCTACCG ATCCGGCAG ATCAGGTGC CCTGCGAAC CCCGCTACCG CAGTCGCG CAGCCGGC CAGCCGGC CAGCCGCC ACCCAGTACC CCGGAACTC CCGGAACTC CCGGAACTC CCGGAACTC CCGGAACCC CCGCACCC CCGGAACCC CCGCGACC CCGCACC CCGCACC CCGCACC CCCC CCTCCC CCTCCC CCCC C	31 CAAGGCCACC TGCCTAGGAT CGCCCGAGT ACATTGTGTT GCTTTGCCAC GCTCTGGGGG GCACAGGGCC AGGTGCCCA CTGACCTGA CTGACCTGA CTGACCTGA CTGACCTGA CTGACCTGA GCCACTGAC GTGACCTGA GCCACTGAC GTGAGCGCA GCCACTGAC GTCAGTGTT TATTTGGCCG AGCAGACCCT AGAGACCT AGAGACCT AGAGACCT AGAGTGCT CCATGGTTCC AGCTGACTCC AGCTGCCGG CCATGGTTCC AGCTGCCGG CCATGGTTCC AGCTGCCGG CATCATTGT CACTTGCTGT GGCGACCCCT AGCTCCCGC AGCTCCCGC CACTTGCTGT CACTTGCTGT AGCTCCCACCCCA CACTCCCCC AGCTCCCCCC CACTTCCCCC CACTTCCCCC AGCTCCCCCC AGCTCCCCCC CACTTCCCCC CACTTCCCCC AGCTCCCCCC AGCTCCCCCC CACTTCCCCC CACTTCCCCCC AGCTCCCCCC AGCTCCCCC CACTTCCCCCC AGCTCCCCCC CACTTCCCCCC AGCTCCCCCC AGCTCCCCCC CACTTCCCCC CACTTCCCCCC AGCTCCCCCC AGCTCCCCCC CACTTCCCCCC AGCTCCCCCC AGCTCCCCCC CACTTCCCCCC CACTTCCCCCC AGCTCCCCCC CACTCCCCCC CACTCCCCC CACTCCCCCC CACCCCCC CACTCCCCCC CACTCCCCC CACTCCCCCC CACTCCCCCC CACTCCCCCC CACTCCCCC CACTCCCCCC CACTCCCCCC CACTCCCCCC CACTCCCCCC CACTCCCCCC CACTCCCCC CACTCCCCCC CACTCCCCCC CACTCCCCCC CACTCCCCCC CACTCCCCCC CACTCCCCC CACTCCCCCC CACTCCCCCC CACTCCCCCC CACTCCCCCC CACTCCCCCC CACTCCCCCC CACTCCCCCC CACTCCCC	CAGAGGGGGA GAGAGCCAG GCAGCCTACAGTACT AGTGCATC AGTGCATC AGTGCATC AGAGTCTG AGGCTGAG GCAGCTGAG GCAGCTGAG GCAGCTGAG CAGCATCTG CAGAGTCAG CAGCATCTG CAGAGTCAG CAGCATCAG GAGGTGAAG CAGCATCAG GAGGTGAG CAGCATCAG GCAGCTCAG GCTCCTGG GCTCCTGGGTAC CAGTGCGGGAC CCGTGGAC CCTTGCAGTGC CCTTGAGAC CCTTGAGAC CCTTGGATAC CCTTGGATCC CCTTGGATGC CCTTGAGTC CCTTGGATGC CCTTGGATGC CCTTGGATGC CCTTGGATGC CCTTGGATGC CCTTGGATCC	GCAAGGGACA CTTCTGGTGG CACAGGGAGA GGCTCCTCAT GGGCGTTGCCT AGCGCATCC CGCGCCATCC AGGCAGGGG CGAGTTGCCT AGGCAGCATC ACCCGACCTC AGGCATCC ACCCGACCTC AGCCATCC ACCCACTC ACCCACTT ATCCTGGCC CGGTGGCCTG CAGGCTGC CGGTTGGAAT GATGTCACC CGGTTGGACT CAGGGGTTG CGGGTTCC CAGGGGTTG CGGGTTGTG ACTCCAGCTG CGGGTTGTG ACTCCC CGGTGCCACT CAGGGGTTCC CAGGGGTTG CCCACTT ACCCCCC CAGTGCCCC CAGTGCCCC CAGGGGTTCCC CAGGGGTTCCC CGGTTGGATTC CCGGTTTGGATTTC CCGGACTCTA	120 180 240 300 360 420 540 660 720 780 840 900 900 1080 1140 1260 1380 1560 1440 1560 1680 1740 1860 1860 1920 1980
50 55 60 65 70 75	Nucleic According sequences of the control of the c	Id Accession lence: 99-8: 11 GGCGCTGGGC TCCTAGCTGA CGCGGGATT CACGGCTCT AGCAATTTC AGCAGTGCA GTTCGGCTGA GTCCCAGGAC CTACAAGGG CTCCAGGAC TCTCCAGGAC TCTCCGGACTT TCCCGGACT TCCCGGACT TCCCGGACA GGCCTGCCAC GGACTGCC GGACTGCC GGACTGCC CGCAACAGC GGAACTGAC CGCAACAGC GGAACTGGCC TGAGGTGCC TCGACTTC CACGGCTGC CCTCTTTC GAGGTGGCC TCGCCTTTC GACTGGCT CCCCGGGAC ACAGGC GCTGGTGCC CCTCGTTCC ACAGGCCGGAG AACAGACTG CCCCCGGGAG AACAGGACTG CACCGGGAG AACAGGACTG CACCGGGAG AACAGGACTG CACCGGGAG AACAGGACTG CACCGGGAG AACAGGACTG CACCGGGAG AACACAGGC CATCACAGGG	# : NM_0000 333 21 TGGGACCTGC CGGCTTTTAC CTGGCAGAGG TACGCCGTG CGGAGGTCC CAGGGTGTC CAGGGTGTC CTGGCCGAC CTGGTGGACA ATCAAGAATG TTCTTCTTCG GTGTGACAG GCACAGGGCC CACTGCCGA CCACTGCCGA CCACTGCCGA ATCAGAGAT TCTTCTTCT GTGTGCACGA GCCACTGCCGA CCACTGCCGA ATCAGGACC CACTGCCGA ATCAGGACC CACTGCCGA ATCAGGACC GCTACCCTGCAA CCCCTGCAA CCCCTGCAA CCCCTGCAA CCCCTGCAA CCCCTGCAA CCCCTGCAA CCCCTGCAA CCCAGTACC CGGAGTCACC CGGAGTCACC CGGAACTC CCGGAAACTC AGGGTGGCCG CTGCAGCCG CTGCAGCCCG CTGCAGCCCG CTGCAGCCCG CTGCAGCCCG CTGCAGCCCG CTGCAGCCCTG CTGCAGCCTG CTGCAGCTG C	31 CAAGGCCACC TGCCTAGGAT CGCCCCGAGT ACATTGTTT GCASCTTTTGCCAC GCTCTGGGGG GCACAGGGGC CTGGTGTCCC CAGCTGCCA TGACCTGA TCAATGACTT CTGCTGGTGG GTGACCTGA CTGACCTGA CTGACCTGA GCACATGAC TGACATGGCG GTCACTGAC GTGACCCCA GGCCACTGAC CTGTGACCCA GCTCACTGA CCAGACCCA AGCAGACCCT TGCCTGAGGG GTTCAGTGT CCGGGGTTCCAGCCG GCATCATTGT CAGCATTCGT CAGCACCCG GCATCATTGT CAGCATTCGT GGGGACCCGT GGGGACCCGT GGGGACCCTA GATCCGCCA GAACCCCTA GAACCCCCA GAACCCCTA	CAGGGGGGA GCAGGGGGGA GCAGCCCAG GCAGCCCAG GCAGCCCAG GCAGCCCAG GCAGCCCAG GCAGCCCAG GCAGCCCAG GCAGCCCCCGCGCCCCCCCGGGCCCCCCCGGGCCCCCCCGGGCCCC	GCAAAGGACA CTTCTGGTGG CACAGGAGA GGCTCCTCAT GTGCTGCCTT AGCGATGACC CGCGCATCC CATGTGGCTG ATCCTGATTA GGGATGACC CAGGCAACA CACAGGAGGGG CGAGTTGCCT AGCCAACTC ACCCGACCTC AGCCAACTC GTGACTGAC CTGATCACT GTGACTGAC CTGACTGCC GTGACTGCC GTGGCCTGC GTGGCCTGC CGGTTGGAC CTGGCCC CCGCCACTT ATCCTGGGCC CAGTTGACCC CAGTTGACCC CAGTTGACCC CAGTTCAGCC CAGGTTGCC CAGGCTCC CAGGCGCC CAGCTCC CAGGCGCC CAGCTCC CAGGCGCC CAGGCTCC CAGGCTCC CAGGCTCC CCGGCTTGC CAGGCTCC CCGGCTTGC CCGGCTTGC CCGGCTTGC CCGGCTTCC CAGGCTCC CCGGCTTCC CCAGCCTCT CCCAGCCTCT CCCAGCCTCT CCCGGCTCC CCGGCTTCC CCGGCTCC CCGGCTCCC CCGCCC CCCC CCCC CCCC CCCC CCCC CCCC CCCC CCCC	120 180 240 360 420 540 660 660 720 780 840 960 1020 1140 1260 1320 1380 1440 1560 1560 1680 1740 1800 1920 1980 2040 2040 2040
505560657075	Nucleic According sequences of the control of the c	Id Accession lence: 99-8: 11 GGGGCTGGGC TCCTAGCTGA CGCCGGGATT CAGCAATTTC AGCCAGTGCA GTTCGCCAGG GTCCCAGGGC TTACAAGGG GTCCCAGGC TTCCCGGACT TCCGGACT TCCGGACT TCCGGACT TCCGGACT GTGGGCGG GTCCCAGGC GCTGCCCAG GCTCCCAGC GCTCCCAGC GCTGGCCAG GCTCGGACAG GCTGGGCC GCTACAGGC GCAACAGC GCAACAGC GCAACAGC CCCACTCC CCCCTCTTCC GACTGGCC CCCCCGGAC GCTGGGCC CCCCCGGGCC ACAGGCC CCCCCGGGCC CCCCGGGAG AACAGACCT CCCCTGGTCC CCCCGGGAG AACAGACCT CCCCTGGTGCC CCCCGGGAG AACAGACCT CCCCTGGTGCC CCCCGGGAG AACAGACCT CACTGCTGCC CCCCGGGAG AACAGGCC CATCACAGGC AACAGGCGAGGAG AACAGGAGGGC AACAGGGGAGGAGGAGGAGGAGGAGGAGGGGCC AACAGGGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGGAGGAGGGG	1 #: NM_0000 333 21 TGGGACCTGC CGGCTTTTAC CTGGCAGAGG TACGCCGCTG CGCAGGTCC CAGGGTGGC CTGGGCCAGACACTC CTGGCCGAC CTGGTGGACA ATCAAGAATG TTCTTCTTCG GTGTGCACGA CCACTGCCGA CCACTGCCGA CCACTGCCGA ATCCAGAGACC CCACTGCCGA ATCCAGATGCC ATCGGGCAGG CTGCTGCTGTG CCTGGCCAGC CGCTCTCTGTTG TGGACCTTC TGGACCTC CCAGCTGCCA CAGCGGCA ACCCCTGCAA CAGCCACCC CAGCGGCA CCCGGAACTC CCGGAACTC CCGGAACTC CCGGAACTC CCGGAACTC AGGGTGGCCT CCTGCAGCCT CCTGCTGCAGCCT CCTGCTGCTGCAGCCT CCTGCTGCTGCAGCCT CCTGCTGCTGCAGCCT CCTGCTGCTGCAGCCT CCTGCTGCAGCCT CCTGCTGCTGCAGCT CCTGCTGCTGCAGCT CCTGCTGCCGC CTGCTGCTGCAGCT CCTGCTGCCGC CTGCTGCTGCCAGCT CCTGCTGCCAGCT CCTGCTGCCAGC CTGCTGCCT CCTGCTGCCAGC CTGCTGCTGCCAGC CTGCTGCTGCCAGC CTGCTGCTGCCAGC CTGCTGCTGCCAGC CTGCTGCTGCCAGC CTGCTGCTGCCAGC CTGCTGCTGCCAGC CTGCTGCCAGC CTGCTGCTGCCAGC CTGCTGCTGCCAGC CTGCTGCTGCCAGC CTGCTGCTGCCAGC CTGCTGCTGCCAGC CTGCTGCTGCCAGC CTGCTGCTGCCAGC CTGCTGCTGCCAGC CTGCTGCTGCCAGC CTGCTGCCTGC CTTGCTGCCAGC CTGCTGCTGCCAC CTGCTGCTGCCAC CTGCTGCTGCCAC CTGCTGCTGCCAC CTGCTGCTGCCAC CTGCTGCTGCCAC CTGCTGCCAC CTGCTGCAC CTGCTGCCAC CTGCTGCCAC CTGCTGCCAC CTGCTGCCAC CTGCTGCCAC CTGCTGCCC CTGCTGCCAC CTGCTGCCAC CTGCTGCCAC CTGCTGCCAC CTGCTGCCAC CTGCTGCCC CTGCTGCCAC CTGCTGCCAC CTGCTGCCAC CTGCTGCCAC CTGCTGCCAC CTGCCAC CTGCTGCCAC CTGCTGCCAC CTGCTGCCAC CTGCTGCCC CTGCTGCCAC CTGCTGCCC CTGCTGCCAC CTGCTGCCC CTGCTGCCAC CTGCTGCCC CTGCCTGC	31 CAAGGCCACC TGCCTAGGAT CGCCCGAGT ACATTGTTT GCAGCTTTCT GCACCTGGGGG GCACAGGGGC CTGGTGCCCA CTGACCTGA TCATGGTGT CTGCTGGTGG CTGTGGTGG CCTGTGGTGG CTGTGGTGG CTGTGACCTGA GCACAGGGCA TGACATGAC TGACAGGCCA TGACAGGCCA TGACAGGCCA TGACATGAC CTGTGAGCGG CCACAGGCCA TGACATGAC TGATGAGCGC AGAAGGTGT TCATTGGCG AGAAGGTGGT CTGAGTGT CTGAGTGC AGAAGTGGT CTGAGTGC CCGTGGTCC AGAAGTGGT CCACTTGCTGG GAGTGCCCA GCATCCCG GGGATCCCG GGGATCCCG GGGACCCTA TCATCGTGT AGTCCACCTA TCATCGTGGC TCACCTAGCCA TCATCGTGGC TCATCCTGG	CAGAGGGGGA GAGGCCAG GCAATCTC GAAGGCTG AGGCTGAAT GCAATCTC CAAGGTCTGAAGGCTGAAGGCTGAAGGCTGAAG CAGACTGAAG CAGACTGAAG CAGACTGAAG CAGACTGAAG CAGACTGAAG CAGACTGAAG CAGACTGAAG CAGACTGAAG CAGACTCAG CAGACTCAG CAGACTCAG CAGCTCCTC CACTGCACCTC CACTGCACCC CTTCGACCC CTCCAGCCC CTCCAGCCC CTCCAGCCC CTCCAGCCC CTCCAGCCC CTCCAGCCC CCCAGCACC CCCCAGGCCC CCCAGGCCC CCCAGCACC CCCAGGCCC CCCAGCACC	GCAAGGGACA CTTCTGGTGG CACAGGGAGA GGCTCCTCAT GTGCTGCCTT AGCGATGACC CGGCCATCC CATGTGGCTG AGCCAGGGGG GGAGTGCCT AGCCACTC AGCCACTGGCC AGTGGCCTG ATCCTGGGCCCACTT ATCCTGGGCCCACTT ATCCTGGGCC CGGTTGGATT CAGGGGTTG CGAGTTCCT CAGGGGTTG CGAGTTCCT CAGGGGTTG ATTCAGGCTG CGAGTTTCCT CAGGGGTTTC CCAGGCTTG AGTGCCATT ATCTGGGCT AGTGCCAGT ATCTAGGCT AGTGCCAGT AGTGCCAGT AGTGCCAGT AGTGCCAGT AGTGCCAGT AGTGCCAGT AGTGCCAGT AGTGCCAGT AGTGCCAGT CCACTGGGCC CCACTCTACC CGGGTTGTGG AGTGCAGTT CCAGGCTTCT CCAGGCTTACC CCACTCTGCCC CCACTCTACC CCACTCTGCCC CCACTCTACC CCACTCTAC CCACTCTGCCC CCACTCTACC CCACTCTAC CCACTCTGCGCC CCACTCTACC CCACTCTAC CCACTCGGCC CCACTCTAC CCACTCGGCC CCACTCTACC CCACTCGGCC CCACTCTACC CCACTCGGCC CCACTCTACC CCACTCGGCC CCACTCTCACC CCACTCGGCC CCACTCTACC CCACTCGGCC CCACTCTCACC CCACTCGGCC CCACTCTCACC CCACTCGGCC CCACTCTCACC CCACTCGGCC CCACTCTCACC CCACTCTCACC CCACTCTCACC CCACTCGGCC CCACTCTCACC CCACTCTCA	120 180 240 300 360 420 660 720 780 1080 1140 1260 1320 1380 1440 1500 1680 1740 1680 1740 1860 1980 2040 2160
50 55 60 65 70 75	Nucleic According sequences of the control of the c	Id Accession lence: 99-8: 11 GGGGCTGGGC TCCTAGCTGA CGCCGGGATT CAGCAATTTC AGCCAGTGCA GTTCGCCAGG GTCCCAGGGC TTACAAGGG GTCCCAGGC TTCCCGGACT TCCGGACT TCCGGACT TCCGGACT TCCGGACT GTGGGCGG GTCCCAGGC GCTGCCCAG GCTCCCAGC GCTCCCAGC GCTGGCCAG GCTCGGACAG GCTGGGCC GCTACAGGC GCAACAGC GCAACAGC GCAACAGC CCCACTCC CCCCTCTTCC GACTGGCC CCCCCGGAC GCTGGGCC CCCCCGGGCC ACAGGCC CCCCCGGGCC CCCCGGGAG AACAGACCT CCCCTGGTCC CCCCGGGAG AACAGACCT CCCCTGGTGCC CCCCGGGAG AACAGACCT CCCCTGGTGCC CCCCGGGAG AACAGACCT CACTGCTGCC CCCCGGGAG AACAGGCC CATCACAGGC AACAGGCGAGGAG AACAGGAGGGC AACAGGGGAGGAGGAGGAGGAGGAGGAGGGGCC AACAGGGGAGGAGGAGGAGGAGGAGGAGGAGGAGGAGGGAGGAGGGG	# : NM_0000 333 21 TGGGACCTGC CGGCTTTTAC CTGGCAGAGG TACGCCGCTG CAGGGTGTGC CAGGGTGTGC CTGGCACACACTC CTGGCACACACTC CTGGCACAC CTGGCCGAC CTGGTGAAC CTGGTGAAC CCACTGCGA CCACTGCGA ATCAAGAATA GCTTCTCTTCG GTGTACACG CCACTGCGA ATCAGAGAC CCACTGCGA ATCAGAGAC CCACTGCGG ATCAGGGC CTTCTGTTG TGGAACTTC GTGAGCCC CATTCTGTTG CCTGGCAG CCAGTCGCC ACCCAGTACC CAGCCGCC ACCCAGTACC CAGGTCAC CAGGTCAC CAGGTCAC CAGGTCAC CAGGTCAC CCGGAACTC CCGCACCC CTGCAGCC CTGCAGCCAC ACTCAGGCCAC ACCCAGCAC ACCCAC AC	31 CAAGGCCACC TGCCTAGGAT CGCCCGAGT ACATTGTGTT GCTTTGCCAC GCTCTGGGGG GCACAGGGCC AGGTGCCCA TGACCTGA TGACCTGA TGACCTGA TGACCTGA GGCACAGGGC GTGAGCGCA GGCACAGGCCA TGACAGGCCA TGACATGAC TGACTGAC TGACTGAC GCACAGCCA TGACAGCCA TGACATGCC AGCAGACCCT AGCAGACCCT AGCAGCCCA AGACAGCCA AGACAGCCA AGACACCGA GCATCATGG CCATGGTCC AGCTGCCG GCATCATGG CACTTGCTG GGGACCCGT AGACCCCA AGACCCCA AGACCCCA AGACCCCA CACCCCA CACCCCA CACCCCC CACTCGCCC CACCCC CACCCC CACCCC CACCCC CCACCCC CCACCC CCACC CCACCC CCACC CCACCC CCACC CCACCC CCACCC CCACCC CCACCC CCACCC CCACCC CCACCC CCACCC CCACCC CCACC CCACCC CCACC CCACC CCACC CC	CAGGGGGGA GCAGCCCAG CCCCCAGGGGC CCCAGGGCCCCCCCC	GCAAGGGACA CTTCTGGTGG CACAGGGAGA GGCTCCTCAT GGGCGTTCA GGCGCATCC CGCGCCATCC AGGCAGGGG CGAGTTGCC AGGCAGCTC AGGCAACCTC AGGCAATCC CGCGACTTC AGCCAATCCT AGCCAATCCT AGCCAATCCT AGCCAATCCT ACCCAGCTG CTGACTGGAC ACCCACTGCC GTGGCCTGC CGGTGGCCTG CCGCCACTT ATCCTGGGCC CGGTTGGACT CAGAGCTCC CGGTTGGACT CAGAGCTCC CGGTTGGACT CAGAGCTCC CGGTTGGACT CCAGGCTG CCGGTTGGACT CCAGGCTG CCGGTTGGACT CCAGACTCT CGGGCTT CCGGTTTGGA ATTGCGTTC CCGGTTTGGATT CCTGGCC CCGGTTGGACT CCGGTTTGGATT CCTGGCC CCGGTTGGACT CCGGTTTGGACT CCGGTTTGGACT CCGGTTCTA CCTGGCCC ACTGGACCC ACCTGGCC ACCTGGACCC ACCTGGACCC ACCTGGACCC ACCTGGACCC ACCTGGACCC ACCTGGACCC ACCTGGACCC ACCTGGACCC ACCTGGACCC	120 180 240 300 360 420 660 720 780 1080 1140 1260 1320 1380 1440 1500 1680 1740 1680 1740 1860 1980 2040 2160

	WO 02	1006112			,		
	WO 02		GCCACGGTGG	CTGAGCTGGA	TGGACTGGAG	CCAGATACTG	2340
	AGTATACGGT	GCATGTGAGG	GCCCATGTGG	CTGGCGTGGA	TGGGCCCCCT	GCCTCTGTGG	2400
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5	CCAGCGACGT	CACTGAACGC	ACCTGGGTAG GGCCCCATGA	GGCACCAGAT	ACTCCCAGGA	AACACAGACTGG	2520 2580
•	CTGCAGAGAT	CCGGGGTCTC	GAAGGTGGAG	TCAGCTACTC	AGTGCGAGTG	ACTGCACTTG	2640
	TCGGGGACCG	CGAGGGCACA	CCTGTCTCCA	TTGTTGTCAC	TACGCCGCCT	GAGGCTCCGC	2700 2760
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10	AGTCCCGGGT	CCTGGGGCCC	GAGCTCAGCA	GCTATCACCT	GGACGGGCTG	GAGCCAGCGA	2880
	CACAGTACCG	CGTGAGGCTG	AGTGTCCTAG	GGCCGGCTGG	AGAAGGGCCC	TCTGCAGAGG	2940 3000
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	CCTGGCGGCC	ACTCAGAGGC	CCTGGCCAGG	AAGTGCCTGG	GTCCCCGCAG	ACACTTCCAG	3120
15	GGATCTCAAG	CTCCCAGCGG	GTGACAGGGC	TAGAGCCTGG	CGTCTCTTAC	ATCTTCTCCC	3180
	TGACGCCTGT	CCTGGATGGT	GTGCGGGGTC GTGGTGTTCC	TACCACATC	CACTCAACACAG	AATGCTCACC	3240 3300
	GTGCGGAGGC	TACGAGGAGG	GTCCTGGAGC	GTCTGGTGTT	GGCACTTGGG	CCTCTTGGGC	3360
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	ACCCAAGIGG	TGGGCGCCGC	CAGCACGTAC	CAGGGGTGAT	GGTTCTGCTA	GTGGATGAAC	3600
	CCTTGAGAGG	TGACATATTC	AGCCCCATCC	GTGAGGCCCA	GGCTTCTGGG	CTTAATGTGG	3660
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65	GCCCCCAGG	CAAGGAGGGC	CCCATCGGCT	TTCCTGGAGA	ACGCGGGCTG	AAGGGCGACC	6120 6180
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	CTGGGGGTGT	GGGAGAGGCA	GGAAGGCCAG	GAGAGAGGGG	AGAACGGGGA	GAGAAAGGAG	6300
	AACGTGGAGA	ACAGGGCAGA	GATGGCCCTC	CTGGACTCCC	TGGAACCCCT	GGGCCCCCCG	6360
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	AGGACGGCAA	CCCGGGTCTA	CCAGGAGAGC	GTGGTATGGC	TGGGCCTGAA	GGGAAGCCGG	6600
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	GACAAGTGGG	GGAGACAGGG	AAGCCGGGAG	CCCCAGGTCG	AGATGGTGCC	AGTGGAAAAG	6900 6960
	ATGGAGACAG	ACCTGGCCCC	GGTGTGCCAG ACGGGGGCCC	CTGGACAGGC	TGTGGTCGGG	CTCCCTGGAG	7020
80	CAAAGGGAGA	GAAGGGAGCC	CCTGGAGGCC	TTGCTGGAGA	CCTGGTGGGT	GAGCCGGGAG	7080
	CCAAAGGTGA	CCGAGGACTG	CCAGGGCCGC	GAGGCGAGAA	GGGTGAAGCT	GGCCGTGCAG	7140
	GGGAGCCCGG	AGACCCTGGG	GAAGATGGTC GTCCCGGGCT	AGAAAGGGGC	TCCAGGACCC	AAAGGTTTCA	7200 7260
_	AGGGAGATCT	GGGCCTCCCT	GGCCTGCCCG	GTGCTCCTGG	TGTTGTTGGG	TTCCCGGGTC	7320
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       AGGATGATGA GTACTCTGAA TACTCCGAGT ATTCTGTGGA GGAGTACCAG GACCCTGAAG
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                                                                              8760
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                                                                              8880
                                                                              8940
25
       ATAATGAGCT GAGATTCAGC ATCCCCTGGA GGAGTCGGGG TCTCAGCAGA ACCCCACTGT
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                                                                              9120
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                                                                              9180
       ACTGGCGTCT GACCCGCCCC TTGACCCAAG CCTGTGATGA CATGGTGCTG ATTCTGGGGG
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       GCATTAAAGC TGCTGTTTTA AAAGGCAAAA AA
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       Protein Accession #: NP 000085.1
35
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       MTLRLLVAAL CAGILAEAPR VRAOHRERVT CTRLYAADIV FLLDGSSSIG RSNFREVRSF
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                                                                               120
       AAILHVADHV FLPQLARPGV PKVCILITDG KSQDLVDTAA QRLKGQGVKL FAVGIKNADP
                                                                               180
40
       EELKRVASQP TSDFFFFVND FSILRTLLPL VSRRVCTTAG GVPVTRPPDD STSAPRDLVL
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                                                                               300
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                                                                               360
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       LRPVII GPTS
                   ILLSWNLVPE ARGYRLEWRR ETGLEPPQKV VLPSDVTRYQ LDGLQPGTEY
                                                                               480
45
       RLTLYTLLEG HEVATPATVV PTGPELPVSP VTDLOATELP GORVRVSWSP VPGATOYRII
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                                                                               600
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                                                                               660
       YOVAVSULRG REEGPAAVIV ARTDPLGPVR TVHVTOASSS SVTITWTRVP GATGYRVSWH
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                   VLRITWVGVT GATAYRLAWG RSEGGPMRHQ ILPGNTDSAE IRGLEGGVSY
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       QPEGGQEQSR VLGPELSSYH LDGLEPATQY
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       ELRVVDTSID
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                                                                             2760
       APGERGEQGR PGPAGPRGEK GEAALTEDDI RGFVRQEMSQ HCACQGQFIA SGSRPLPSYA
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       DEGSCTAYTL RWYHRAVTGS TEACHPFVYG GCGGNANRFG TREACERRCP PRVVQSQGTG
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WO 02/086443

Seq ID NO: 64 DNA sequence Nucleic Acid Accession #: NM_006945 Coding sequence: 1-219

•	Coding sequence: 1-219						
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10			TCAGCAGTGC TCCACCGAAG		ATCCTCCTGT	GACACCTTCC	180
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	PPCOPKYPPK MSYQQQQCKQ		PKCPEPCPPP	KCPEPCPPPK	CPQPCPPQQC	QQKYPPVTPS	60
20	Nucleic Act	66 DNA sequid Accession lence: 639-2	ı#: №М_005	6629.1			
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	CCGCCGCCGG	GAAGGAGAGG	GCGAGGCGCG	CCCGAGCCGC	CGCCGCCGCC	GCCACCGCCG	120
			GGAGTCGCGG				180 240
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			GCGCGCCCC				420
			GGACTGCTTC				480 540
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			GTGCGGCCCG				660
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			TGGACGCGCC				840
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			GTCCTGATCG TTCATGAAGG				960 1020
			TACGCCTCCA				1080
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45			ACCTGGAACA				1200 1260
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	CAGGGGCCCT	CAACTGGGAG	GTGACCCTTT	GTCTGCTGGC	CTGCTGGGTG	CTGGTCTACT	1380
50			AAATCCACGG				1440
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			AACAACTGCT				1680
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			GCCTCCTACT				1980 2040
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						CCAGGCTCTG	2880
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	CCTACCTTAC	CCCTCTGCCC	CTAGCCAAGG TGTTGCGTGT	AGTGTGAATT	TATAGATCTA	ACTTTCATÁG	3420
	GTCCCCAGCC	CCAGACTGGA	TTTGGAAAAGT	GCATGGTGGG	GGCCTCGGGG	CTGTCCCCAC	3480 3540
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	GGGCTGCTAA	CCTGGCCTGC	TCAGGCTTCC	CACCCTGTGC	GGGGCACACC	CCCAGGAAGG	3660
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	GSPQVWIDAG SILGFMAAEQ GVEGFITGLL TTLLWQAFWE	TQIFFSYAIG GVHISKVAES DLLPASYYFR CVVVAWVYGA	VYFTATFPYV LGALTALGSY GPGLAFIAYP FQREISVALC DRFMDDIACM	NRFNNNCYKD RAVTLMPVAP CALCFVIDLS IGYRPCPWMK	AIILALINSG LWAALFFFML MVTDGGMYVF WCWSFFTPLV	TSFFAGFVVF LLLGLDSQFV QLFDYYSASG CMGIFIFNVV	300 360 420 480 540
20			GWAFALSSML PVSESSKVVV		LRAKGTMAER	MÖHLIÐÞIMG	600
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55			CACCATGCCC				420
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			CATCAGCTGT				540 600
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			CTGGCAGAAC				900
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			GACCATTCAC				1140
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	JACCHMANA	AMMANAAAA	· maran)				

Seq ID NO: 69 Protein sequence: Protein Accession #: NP_068772.1

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25	-						
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Seq ID NO: 71 Protein sequence: Protein Accession #: AAH06529.1

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		AATTCATCCT					540
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		CATCAGCGTC					900
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		TGAACCAGTC					1380
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• •		AGGAGGGGAT					1620
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		GTACTTCCCG					2040
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Seq ID NO: 73 Protein sequence: Protein Accession #: AAC51128.1

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55			ATTGAAGCAA				7140
55							7200
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			GACAGAGCTG				7320
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60	ATGATGCCAA	AGTTAAAACC	AGTAGAACTC	CGAGAACTTC	TGAACCCCGT	TGTGGAATTC	7440
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80							8640
JV	CGTTTTCTTA	ATACCACCTT	CTCTTTCTTT	CONCELLING	CTCCCTCCTAT	TOTOGRUAL T	
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	GCCAGCCTAC	AGCAGCCCGT	GGGCATCCGC	CTGCTAGAGG	AGGCTCTGCT	CCGCCTGCTG	8760
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                                                                                         240
                                                                                         300
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VYEMFRKDDP RLSFTRQSFV DRSLLTLLWH CSLDALREFF STIVVDAIDV LKSRFTKLNE
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                                                                                     1860
                                                                                     1920
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         QEEKOKCYND LLASAKKOLE VEROTITOLS FELSEFRRKY EETOKEVHNL NOLLYSORRA
DVOKLEDDRH KTEKIOKLRE ENDIARGKLE EEKKRSEELL SOVOSLYTSL LKOOEEOTRV
                                                                                                 120
                                                                                                 180
20
         ALLEQOMOAC TLDPENEKLD ROHVOHOLHV ILKELRKARK NNTVGILETA S
         Seg ID NO: 145 DNA sequence
         Nucleic Acid Accession #: NM_001168
         Coding sequence: 50..478
25
                                      21
         CCGCCAGATT TGAATCGCGG GACCCGTTGG CAGAGGTGGC GGCGGCGGCA TGGGTGCCCC
                                                                                                   60
30
         GACGTTGCCC CCTGCCTGGC AGCCCTTTCT CAAGGACCAC CGCATCTCTA CATTCAAGAA
          CTGGCCCTTC TTGGAGGGCT GCGCCTGCAC CCCGGAGCGG ATGGCCGAGG CTGGCTTCAT
                                                                                                 180
         CCACTGCCC ACTGAGAACG AGCCAGACTT GGCCCAGTGT TTCTTCTGCT TCAAGGAGCT
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                                                                                                 240
                                                                                                 300
         CGCTTTCCTT TCTGTCAAGA AGCAGTTTGA AGAATTAACC CTTGGTGAAT TTTTGAAACT
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35
                                                                                                 420
                                                                                                 480
          CCTCTGGCCG GAGCTGCCTG GTCCCAGAGT GGCTGCACCA CTTCCAGGGT TTATTCCCTG
         GTGCCACCAG CCTTCCTGTG GGCCCCTTAG CAATGTCTTA GGAAAGGAGA TCAACATTTT
                                                                                                  600
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                                                                                                 660
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GGGGGCTCAT TTTTGCTGTT TTGATTCCCG GGCTTACCAG GTGAGAAGTG AGGGAGGAAG
40
                                                                                                 780
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                                                                                                 840
         GTGAATGTGT CTGGACCTCA TGTTGTTGAG GCTGTCACAG TCCTGAGTGT GGACTTGGCA
GGTGCCTGTT GAATCTGAGC TGCAGGTTCC TTATCTGTCA CACCTGTGCC TCCTCAGAGG
                                                                                                  900
                                                                                                 960
45
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         GTGATGAGAG AATGGAGACA GAGTCCCTGG CTCCTCTACT GTTTAACAAC ATGGCTTTCT
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                                                                                                1080
                                                                                                1140
         AGCCATTCTA AGTCATTGGG GAAACGGGGT GAACTTCAGG TGGATGAGGA GACAGAATAG
                                                                                                1200
         AGTGATAGGA AGCGTCTGGC AGATACTCCT TTTGCCACTG CTGTGTGATT AGACAGGCCC AGTGAGCCGC GGGGCACATG CTGGCCGCTC CTCCCTCAGA AAAAGGCAGT GGCCTAAATC
                                                                                                1260
50
                                                                                                1320
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                                                                                                1380
         TCTGTCAGCC CAACCTTCAC ATCTGTCACG TTCTCCACAC GGGGGAGAGA CGCAGTCCGC CCAGGTCCCC GCTTTCTTTG GAGGCAGCAG CTCCCGCAGG GCTGAAGTCT GGCGTAAGAT
                                                                                                1440
         GATGGATTTG ATTCGCCCTC CTCCCTGTCA TAGAGCTGCA GGGTGGATTG TTACAGCTTC
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         MGAPTLPPAW QPFLKDHRIS TFKNWPFLEG CACTPERMAE AGFIHCPTEN EPDLAQCFFC
         PKELEGWEPD DDPIEEHKKH SSGCAFLSVK KQFEELTLGE FLKLDRERAK NKIAKETNNK
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65
         KKEFEETAKK VRRAIEOLAA MD
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70
                                                                                 51 '
         GCGCGCAGCG CTGGTACCCC GTTGGTCCGC GCGTTGCTGC GTTGTGAGGG GTGTCAGCTC
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                                                                                                 120
75
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                                                                                                 360
                                                                                                 420
80
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         AACCCTGATG ACCCGCTCAT GGCTGACATA TCCTCAGAAT TTAAATATAA TAAGCCAGCC
TTCCTCAAGA ATGCCAGACA GTGGACAGAG AAGCATGCAA GACAGAAACA AAAGGCTGAT
                                                                                                 540
                                                                                                 600
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         CAGAAAAGGA AGGCCAGTCA GCTAGTAGGC ATAGAAAAGA AATTTCATCC TGATGTTTAG
GGGACTTGTC CTGGTTCATC TTAGTTAATG TGTTCTTTGC CAAGGTGATC TAAGTTGCCT
                                                                                                 720
85
                                                                                                 780
         ACCTTGAATT, TTTTTTTAAA TATATTTGAT GACATAATTT TTGTGTAGTT TATTTATCTT
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Sec ID NO: 148 Protein sequence: Protein Accession #: NP 054895.1 5 51 MQRASRLKRE LHMLATEPPP GITCWQDKDQ MDDLRAQILG GANTPYEKGV FKLEVIIPER YPPEPPQIRF LTPIYHPNID SAGRICLDVL KLPPKGAWRP SLNIATVLTS IQLLMSEPNP 120 10 DDPLMADISS EFKYNKPAFL KNARQWTEKH ARQKQKADEE EMLDNLPEAG DSRVHNSTQK RKASOLVGIE KKFHPDV Seq ID NO: 149 DNA sequence Nucleic Acid Accession #: NM_003812 15 Coding sequence: 224-2722 51 31 TCCTCTGCGT CCCGCCCCGG GAGTGGCTGC GAGGCTAGGC GAGCCGGGAA AGGGGGCGCC 20 GCCCAGCCC GAGCCCCGCG CCCCGTGCCC CGAGCCCGGA GCCCCCTGCC CGCGGCGGCA 120 CCATGCGCGC CGAGCCGGCG TGACCGGCTC CGCCCGGGC CGCCCGCAG CTAGCCCGGC 180 GCTCTCGCCG GCCACACGGA GCGGCGCCCG GGAGCTATGA GCCATGAAGC CGCCCGGCAG 240 CAGCTCGCGG CAGCCGCCCC TGGCGGGCTG CAGCCTTGCC GGCGCTTCCT GCGGCCCCCA ACGCGGCCC GCCGGCTCGG TGCCTGCCAG CGCCCCGGCC CGCACGCCGC CCTGCCGCCT 360 25 GCTTCTCGTC CTTCTCCTGC TGCCTCCGCT CGCCGCCTCG TCCCGGCCCC GCGCCTGGGG 420 GGCTGCTGCG CCCAGCGCTC CGCATTGGAA TGAAACTGCA GAAAAAAATT TGGGAGTCCT 480 GGCAGATGAA GACAATACAT TGCAACAGAA TAGCAGCAGT AATATCAGTT ACAGCAATGC AATGCAGAAA GAAATCACAC TGCCTTCAAG ACTCATATAT TACATCAACC AAGACTCGGA 600 AAGCCCTTAT CACGTTCTTG ACACAAAGGC AAGACACCAG CAAAAACATA ATAAGGCTGT CCATCTGGCC CAGGCAAGCT TCCAGATTGA AGCCTTCGGC TCCAAATTCA TTCTTGACCT 660 30 CATACTGAAC AATGGTTTGT TGTCTTCTGA TTATGTGGAG ATTCACTACG AAAATGGGAA 780 ACCACAGTAC TCTAAGGGTG GAGAGCACTG TTACTACCAT GGAAGCATCA GAGGCGTCAA 840 AGACTCCAAG GTGGCTCTGT CAACCTGCAA TGGACTTCAT GGCATGTTTG AAGATGATAC 900 CTTCGTGTAT ATGATAGAGC CACTAGAGCT GGTTCATGAT GAGAAAAGCA CAGGTCGACC 960 35 ACATATAATC CAGAAAACCT TGGCAGGACA GTATTCTAAG CAAATGAAGA ATCTCACTAT GGAAAGAGGT GACCAGTGGC CCTTTCTCTC TGAATTACAG TGGTTGAAAA GAAGGAAGAG 1020 1080 AGCAGTGAAT CCATCACGTG GTATATTTGA AGAAATGAAA TATTTGGAAC TAATGATCAC AAAACGTATA AGAAGCATCG CTCTTCTCAT GCACATACCA ACAACTTTGC AAAGTCCGTG GTCAACCTTG TGGATTCTAT TTACAAGGAG CAGCTCAACA CCAGGGTTGT 1200 1260 40 CCTGGTGGCT GTAGAGACCT GGACTGAGAA GGATCAGATT GACATCACCA CCAACCCTGT GCAGATGCTC CATGAGTTCT CAAAATACCG GCAGCGCATT AAGCAGCATG CTGATGCTGT 1380 GCACCTCATC TCGCGGGTGA CATTTCACTA TAAGAGAAGC AGTCTGAGTT ACTTTGGAGG 1440 TOTCTGTTCT CGCACAGAG GAGTTGGTGT GAATGAGTAT GGTCTTCCAA TGGCAGTGGC 1500 ACAAGTATTA TCGCAGAGCC TGGCTCAAAA CCTTGGAATC CAATGGGAAC CTTCTAGCAG 1560 45 AAAGCCAAAA TGTGACTGCA CAGAATCCTG GGGTGGCTGC ATCATGGAGG AAACAGGGGT GTCCCATTCT CGAAAATTTT CAAAGTGCAG CATTTTGGAG TATAGAGACT TTTTACAGAG 1620 1680 1740 AGGAGGTGGA GCCTGCCTTT TCAACAGGCC AACAAAGCTA TTTGAGCCCA CGGAATGTGG AAATGGATAC GTGGAAGCTG GGGAGGAGTG TGATTGTGGT TTTCATGTGG AATGCTATGG 1800 ATTATGCTGT AAGAANTGTT CCCTCTCCAA CGGGGCTCAC TGCAGCGACG GGCCCTGCTG
TAACAATACC TCATGTCTTT TTCAGCCACG AGGGTATGAA TGCCGGGATG CTGTGAACGA 1860 50 1920 GTGTGATATT ACTGAATATT GTACTGGAGA CTCTGGTCAG TGCCCACCAA ATCTTCATAA 1980 GCAAGACGGA TATGCATGCA ATCAAAATCA GGGCCGCTGC TACAATGGCG AGTGCAAGAC CAGAGACAAC CAGTGTCAGT ACATCTGGGG AACAAAGGCT GCAGGGTCTG ACAAGTTCTG 2040 2100 CTATGAAAAG CTGAATACAG AAGGCACTGA GAAGGGAAAC TGCGGGAAGG ATGGAGACCG 2160 55 GTGGATTCAG TGCAGCAAAC ATGATGTGTT CTGTGGATTC TTACTCTGTA CCAATCTTAC 2220 TCGAGCTCCA CGTATTGGTC AACTTCAGGG TGAGATCATT CCAACTTCCT TCTACCATCA 2280 AGGCCGGGTG ATTGACTGCA GTGGTGCCCA TGTAGTTTTA GATGATGATA CGGATGTGGG 2340 CTATGTAGAA GATGGAACGC CATGTGGCCC GTCTATGATG TGTTTAGATC GGAAGTGCCT 2400 ACAAATTCAA GCCCTAAATA TGAGCAGCTG TCCACTCGAT TCCAAGGGTA AAGTCTGTTC 2460 60 GGGCCATGGG GTGTGTAGTA ATGAAGCCAC CTGCATTTGT GATTTCACCT GGGCAGGGAC 2520 AGATTGCAGT ATCCGGGATC CAGTTAGGAA CCTTCACCCC CCCAAGGATG AAGGACCCAA 2580 GGGTCCTAGT GCCACCAATC TCATAATAGG CTCCATCGCT GGTGCCATCC TGGTAGCAGC 2640 TATTGTCCTT GGGGGCACAG GCTGGGGATT TAAAAATGTC AAGAAGAGAA GGTTCGATCC 2700 TACTCAGCAA GGCCCCATCT GAATCAGCTG CGCTGGATGG ACACCGCCTT GCACTGTTGG 2760 65 ATTCTGGGTA TGACATACTC GCAGCAGTGT TACTGGAACT ATTAAGTTTG TAAACAAAAC 2820 CTTTGGGTGG TAATGACTAC GGAGCTAAAG TTGGGGTGAC AAGGATGGGG TAAAAGAAAA 2880 CTGTCTCTTT TGGAAATAAT GTCAAAGAAC ACCTTTCACC ACCTGTCAGT AAACGGGGGA GGGGGCAAAA GACCATGCTA TAAAAAGAAC TGTTCCAGAA TCTTTTTTT TCCCTAATGG 3000 ACGAAGGAAC AACACACAC CAAAAATTAA ATGCAATAAA GGAATCATTA AAAA 70 Seq ID NO: 150 Protein sequence: Protein Accession #: NP 003803 75 41 31 51 MKPPG99SRQ PPLAGCSLAG ASCGPQRGPA GSVPASAPAR TPPCRLLLVL LLLPPLAASS RPRAWGAAAP SAPHWNETAE KNLGVLADED NTLQQNSSSN ISYSNAMQKE ITLPSRLIYY 120 INODSESPYH VLDTKARHOO KHNKAVHLAQ ASFQIEAFGS KFILDLILNN GLLSSDYVEI 180 80 HYENGKPQYS KGGEHCYYHG SIRGVKDSKV ALSTCNGLHG MFEDDTFVYM IEPLELVHDE KSTGRPHIIQ KTLAGQYSKQ MKNLTMERGD QWPFLSELQW LKRRKRAVNP SRGIFEEMKY LELMIVNDHK TYKKHRSSHA HTNNFAKSVV NLVDSIYKEQ LNTRVVLVAV ETWTEKDQID ITTNPVQMLH EFSKYRQRIK QHADAVHLIS RVTFHYKRSS LSYFGGVCSR TRGVGVNEYG 300 360 420 LPMAVAQVLS QSLAQNLGIQ WEPSSRKPKC DCTESWGGCI MEETGVSHSR KFSKCSILEY 480 85 RDFLORGGGA CLENRPTKLF BPTECGNGYV EAGEECDCGF HVECYGLCCK KCSLSNGAHC 54D SDGPCCNNTS CLFQPRGYEC RDAVNECDIT EYCTGDSGQC PPNLHKQDGY ACNQNQGRCY 600 NGECKTRDNQ CQYIWGTKAA GSDKFCYEKL NTEGTEKGNC GKDGDRWIQC SKHDVFCGFL

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	LDRKCLQIQA	IGQLQGEIIP LNMSSCPLDS TNLIIGSIAG	KGKVCSGHGV	CSNEATCICD	FTWAGTDCSI	RDPVRNLHPP	720 780
5	Nucleic Aci	151 DNA sec d Accession lence: 250-1	#: NM_0239	15			
10	1	11	21	31	41	51	
10	 	TTTCGTTTTC	ATGCTTTACC	AGAAAATCCA	CTTCCCTGCC	GACCTTAGTT	60
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		AGCCAGCCAC					180 240
15	AACTGAAGAA	AATCGTCCCC TGGGGTTCAA	CTTGACGCTT	GCAAAATTAC	CAAATAACGA	GCTGCACGGC	300
	CAAGAGAGTC	ACAATTCAGG	CAACAGGAGC	GACGGGCCAG	GAAAGAACAC	CACCCTTCAC	360
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	TTGCTGAATG	AAAACATAGT	GGTTGCAGAC	CTCATAATGA	CGCTGACATT	TCCATTTCGA	540
20	ATAGTCCATG	ATGCAGGATT	TGGACCTTGG	TACTTCAAGT	TTATTCTCTG	CAGATACACT	600
	TCAGTTTTGT	TTTATGCAAA TGAAGGTGGT	CATGTATACT	TCCATCGTGT	TCCTTGGGCT	CATAAGCATT	660 720
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		TTAAAAAAAA		AIAIGIACAA	AGIGIAMIA	<i>A</i>	-500
35							
	Sea ID NO:	152 Protei	a sequence:				
		cession #: 1					
40	į	11	21	31	41	51	
	MGENT.TLAKI.	PNNELHGQES	HNSGNRSDGP	GKNTTLHNEF	DTIVLPVLYL	 IIFVASILLN	60
	POPMENTEN	z mnomrodno				DET COMMOUNT	
	GLAVWIFFHI	RNKTSFIFYL	KNIVVADLIM	TLTFPFRIVH	DAGFGPWYFK	FILCRITSVL	120
45	FYANMYTSIV	FLGLISIDRY	LKVVKPFGDS	RMYSITFTKV	LSVCVWVIMA	VLSLPNIILT	180
45	FYANMYTSIV NGQPTEDNIH ISQSSRKRKH	FLGLISIDRY DCSKLKSPLG NQSIRVVVAV	LKVVKPFGDS VKWHTAVTYV FFTCFLPYHL	RMYSITFTKV NSCLFVAVLV CRIPFTFSHL	LSVCVWVIMA ILIGCYIAIS DRLLDESAQK	VLSLPNIILT RYIHKSSRQF ILYYCKEITL	
45	FYANMYTSIV NGQPTEDNIH ISQSSRKRKH	FLGLISIDRY	LKVVKPFGDS VKWHTAVTYV FFTCFLPYHL	RMYSITFTKV NSCLFVAVLV CRIPFTFSHL	LSVCVWVIMA ILIGCYIAIS DRLLDESAQK	VLSLPNIILT RYIHKSSRQF ILYYCKEITL	180 240
	FYANMYTSIV NGQPTEDNIH ISQSSRKRKH FLSACNVCLD Seq ID NO:	FLGLISIDRY DCSKLKSPLG NQSIRVVVAV PIIYFFMCRS 153 DNA sec	LKVVKPFGDS VKWHTAVTYV PFTCFLPYHL FSRRLFKKSN Quence	RMYSITFTKV NSCLFVAVLV CRIPFTFSHL IRTRSESIRS	LSVCVWVIMA ILIGCYIAIS DRLLDESAQK	VLSLPNIILT RYIHKSSRQF ILYYCKEITL	180 240
45 50	FYANMYTSIV NGQPTEDNIH ISQSSRKRKH FLSACNVCLD Seq ID NO: Nucleic Ac:	FLGLISIDRY DCSKLKSPLG NQSIRVVVAV PIIYFFMCRS 153 DNA see id Accession	LKVVKPFGDS VKWHTAVTYV PFTCFLPYHL FSRRLFKKSN Quence 1 #: D80008	RMYSITFTKV NSCLFVAVLV CRIPFTFSHL IRTRSESIRS	LSVCVWVIMA ILIGCYIAIS DRLLDESAQK	VLSLPNIILT RYIHKSSRQF ILYYCKEITL	180 240
	FYANMYTSIV NGQPTEDNIH ISQSSRKRKH FLSACNVCLD Seq ID NO: Nucleic Ac: Coding sequ	FLGLISIDRY DCSKLKSPLG NQSIRVVVAV PIIYFFMCRS 153 DNA sec id Accession Lence: 149-	LKVVKPFGDS VKWHTAVTYV FFTCFLPYHL FSRRLFKKSN Quence 1 #: D80008	RMYSITFTKV NSCLFVAVLV CRIPFTFSHL IRTRSESIRS	LSVCVWVIMA ILIGCYIAIS DRLLDESAQK LQSVRRSEVR	VLSLPNIILT RYIHKSSRQF ILYYCKEITL IYYDYTDV	180 240
50	FYANMYTSIV NGQPTEDNIH ISQSSRKRKH FLSACNVCLD Seq ID NO: Nucleic Ac:	FLGLISIDRY DCSKLKSPLG NQSIRVVVAV PIIYFFMCRS 153 DNA see id Accession	LKVVKPFGDS VKWHTAVTYV PFTCFLPYHL FSRRLFKKSN Quence 1 #: D80008	RMYSITFTKV NSCLFVAVLV CRIPFTFSHL IRTRSESIRS	LSVCVWVIMA ILIGCYIAIS DRLLDESAQK	VLSLPNIILT RYIHKSSRQF ILYYCKEITL	180 240
	FYANMYTSIV NGQPTEDNIH ISQSSRKRKH FLSACNVCLD Seq ID NO: Nucleic Ac: Coding sequ 1	FLGLISIDRY DCSKLKSPLG NQSIRVVVAV PILYFFMCRS 153 DNA see id Accession dence: 149- 11 AAAGCGCGGA	LKVVKPFGDS VKWHTAVTYV FFTCFLPYHL FSRRLFKKSN QUENCE 1 #: D80008 739 21] GCGGAGGCCG	RMYSITFTKV NSCLFVAVLV CRIPFTFSHL IRTRSESIRS 1.1 31 1 AGGCGAGAGC	LSVCVWVIMA ILIGCYIAIS DRLLDESAQK LQSVRRSEVR 41 CTGGCGCTGT	VLSLPNIILT RYIHKSSRQF ILYYCKEITL IYYDYTDV 51 AGGACTAGAA	180 240 300
50	FYANMYTSIV NGQPTEDNIH ISGSRKRKH PLSACNVCLD Seq ID NO: Nucleic Ac: Coding sequ 1 GTTCGGCGCC CGAAAGGAGT	FLGLISIDRY DCSKLKSPLG NQSIRVVVAV PITYFFMCRS 153 DNA see id Accession nence: 149- 11 AAAGCGCGGA GAGGCGCCGA	LKVVKPFGDS VKWHTAVTYV FFTCFLPYHL FSRRLFKKSN Quence 1 #: D80008 739 21 GCGGAGGCCG GAGCCCAGAT	RMYSITFTKV NSCLFVAVLV CRIPFTFSHL IRTRSESIRS 1.1 31 AGGCGAGAGA ACCATTTGG	LSYCVWVIMA ILIGCYIAIS DRLLDESAQK LQSVRRSEVR 41) CTGGCGCTGT CGTGAGAGCT	VLSLPNIILT RYIHKSSRQF ILYYCKETTL IYYDYTDV 51 AGGACTAGAA GGTGGTTGGC	180 240 300
50	FYANMYTSIV NGQPTEDNIH ISQSSRKRKH PLSACNVCLD Seq ID NO: Nucleic Ac: Coding sequ GTTCGGCGCC CGAAAGGAGT AAGGCCGCGG CGAACGAGT CGASCTGCAT	FLGLISIDRY DCSKLKSPLG NQSIRVVVAV PITYFFMCRS 153 DNA see id Accession tence: 149-' 11 AAAGCGCGGGA GAGGGCCCGA GAGGGCCCGGA GGGGGCCCGG	LKVVKPFGDS VKWHTAVTYV FFTCFLPYHL FSRRLFKKSN Quence 1 #: D80006 739 21 GCGGAGGCCG GAGCCCAGAT CGTCCGCAT AAGGGCAACT	RMYSITFTKV NSCLFVAVLV CRIPFTFSHL IRTRSESIRS 1.1 31 AGGCGAGAGG ACCATTTGG GTTCTGCGAA GCCTGCCTTC	LSYCVWVIMA ILIGCYIAIS DRLLDESAQK LQSYRRSEVR 41) CTGGCGCTGT CGTGAGAGCT AAAGCCATGG AACGAGGATG	VLSLPNIILT RYIHKSSROF ILYYCKEITL IYYDYTDV 51 AGGACTAGAA GGTGGTTGGC AACTGATCCG GACTCAGACA	190 240 300 60 120 180 240
50	FYANMYTSIV NGQPTEDNIH ISQSSRKRKH PLSACNVCLD Seq ID NO: Nucleic Ac: Coding sequ 1 GTTCGGCGCC CGAAAGGAGT AAGGCCGCAA AGTTCTGGAA AGTTCTGGAG	FLGLISIDRY DCSKLKSPLG NQSIRVVVAV PITYFFMCRS 153 DNA see id Accession nence: 149- 11 AAAGCGCGGA GAGTGGAAG GAGGGCCCG GAGATGAAAG GAGGAGCAAG GAGATGAAAG	LKVVKPFGDS VKWHTAVTYV FFTCFLPYHL FSRRLFKKSN Quence 1 #: D80008 739 21 GCGGAGGCCG GAGCCCAGAT CGTCGCCAT AAGGGCAACT CTTTGTATGA	RMYSITFTKV NSCLFVAVLV CRIPFTFSHL IRTRSESIRS .1 31 AGGCGAGAGC ACCATTTTGG GTTCTGCGAA GCCTGCCTTC ACAAAACCAG	LSYCVWVIMA ILIGCYIAIS DRLLDESAQK LQSVRRSEVR 41) CTGGCGCTGT CGTGAGAGCT AAAGCCATGG AACGAGGATG TCTGATGTGA	VLSLPNIILT RYIHKSSRQF ILYYCKETTL IYYDYTDV 51 AGGACTAGAA GGTGGTTGGC AACTGATCCG GACTCAGACA ATGAAGCAAA	180 240 300 60 120 180 240 300
50	FYANMYTSIV NGQPTEDNIH ISQSSRKRKH PLSACNVCLD Seq ID NO: Nucleic Ac: Coding sequ [FLGLISIDRY DCSKLKSPLG NQSIRVVVAV PIIYFFMCRS 153 DNA see id Accession ence: 149-' 11 AAAGCGCGGA GAGTGGAAG GAGTGGAAG GCGCCCG GAGATGAAAG CCAAGTGGTAT TGCACTGTAG	LKUVKPFGDS VKWHTAVTYV FFTCFLPYHL FFRRLFKKSN Quence 1 #: D80008 739 21 GCGGAGGCCG GAGCCCAGAT CGTCGCAAT CGTCGCAAT CGTTCGTATGA TGATACCAAC CATACCTGTA	RMYSITFTKV NSCLFVAVLV CRIPFTFSHL IRTRSESIRS 1.1 31 AGGCGAGAGG ACCATTTTGG GTTCTGCGAA GCCTGCCTTC ACAAAACCAG TATCAAATTT TGACCGCTTG	LSYCVWYIMA ILIGCYIAIS DRLLDESAQK LQSYRRSEVR 41 CTGGCGCTGT CGTGAGAGCT AAAGCCATGG AACGAGGATG TCTGATGTGACCGTCT TCTGGTGATCA	VLSLPNIILT RYIHKSSRQF ILYYCKEITL IYYDYTDV 51 AGGACTAGAA GGTGGTTGGG GACTCAGACA ATGAGCCAA ATGAGCAAA CTCTGTTAGAG GAGCACTCAG	180 240 300 60 120 180 240 300 360 420
50	FYANMYTSIV NGQPTEDNIH ISGSRKRKH PLSACNVCLD Seq ID NO: Nucleic Ac: Coding sequ 1 GTTCGGCGCC CGAAAGGAGT AAGGCCGCGG CGAACTGGCAT AGGTCTGGAG GTCAGGTGGA AAATCGACGC ATGGGAAATAT	FLGLISIDRY DCSKLKSPLG NQSIRVVVAV PITYFFMCRS 153 DNA see id Accession ence: 149-' 11 AAAGCGCGGA GAGTGGAAG GCGCGCCG GAGTGAAAG CGAGTGAAAG CGAAGTGATT TGCACTGTAG GGTAGGGTCT	LKVVKPFGDS VKWHTAVTYV FFTCFLPYHL FFRRLFKKSN Quence 1 #: D80006 739 21 GCGGAGGCCG GAGCCCAGAT CGTCGCCAT AAGGGCAACT CTTTGTATGA TGATACCTATA TGCCAAATGC	RMYSITFTKV NSCLFVAVLV CRIPFTFSHL IRTRSESIRS .1 31 AGGCGAGAGC ACCATTTTGG GTTCTGCGAA GCCTGCCTTC ACAAAACCAG TATCAAATT TGACCGCTTG ATTACGATTT TGACGATTT	LSYCWYIMA ILIGCYIAIS DRLLDESAQK LQSYRRSEVR 41) CTGGCGCTGT CGTGAGAGCT AAAGCCATGG TCTGAGTGAG TCTGATGTGA CGACACTGTT CTTCGGATCA CACATGGCTG	VISLPNIILT RYIHKSSRQF ILYYCKETTL IYYDYTDV 51 AGGACTAGAA GCTGGTTGGC AACTGATCCG GACTCAGACA ATGAAGCAAA CTCTGTTAAG GAGCACTCAG CTGAAGAAAA	180 240 300 60 120 180 240 300 360 420 480
50 55 60	FYANMYTSIV NGQPTEDNIH ISGSSRKRKH PLSACNVCLD Seq ID NO: Nucleic Ac: Coding sequ 1 GTTCGGCGCC CGAAAGGAGT AAGGCCGCGG CGAGCTGCAT AGTTCTGGAG GTCAGGTGGAA AAATCGACGC ATGGGAATAT GGAGTGGTTTT	FLGLISIDRY DCSKLKSPLG NQSIRVVVAV PITYFFMCRS 153 DNA see id Accession sence: 149- 11 AAAGCGCGGA GAGTGGGAAG CGCGCCCG GAGTGGGAAG CGAGTGATT TGCACTGTAG GGTAGCGTCT AATAATTATA	LKVVKPFGDS VKWHTAVTYV FFTCFLPYHL FFRRLFKKSN Quence 1 #: D80008 739 21] GCGGAGGCCG GAGCCCAGAT CGTCCGCCAT AAGGGCAACT CTTTGTATGA TGATACCTAC TGATACCTGTA TGCCAAATGC AAAGATCTCT AAAGATCTCT	RMYSITFTKV NSCLFVAVLV CRIPFTFSHL IRTRSESIRS 1.1 31 AGGCGAGAGC GTTCTGCGAA GCTTGCGTTG ACAAAACCAG TATCAAATTT TGACACGTTG TTTCACGTTT TGCTACTTAT	LSYCWYIMA ILIGCYIAIS DRLLDESAQK LQSYRRSEVR 41 CTGGCGCTGT CGTGAGAGCT AAAGCCATGG AACGAGATT CTTCGGATCA CGACACTGTT CTTCGGATCA ACATGCTG ATGAGGTCA ATGAGGTCA	VISLPNIILT RYIHKSSROF ILYYCKETTL IYYDYTDV 51 AGGACTAGAA GGTGGTTGGC AACTGATCCG GACTCAGACA ATGAAGCAAA CTCTGTTAAG GAGCACTCAG CTGAAGAAAT TGGGAGGAGA	180 240 300 60 120 180 240 360 420 480 540
50	FYANMYTSIV NGQPTEDNIH ISQSSRKRKH FLSACNVCLD Seq ID NO: Nucleic Ac: Coding sequ (GTTCGGCGCC CGAAAGGAGT AAGGCTGCAG AAGGCTGCAG GTCAGGTGGA AAAATCGACGC ATGGGAATAT GGAGTGGTTT TGAAGGTTTT TGAAGGTTT TGAAGGTTTAAAA	FLGLISIDRY DCSKLKSPLG NQSIRVVVAV PIIYFFMCRS 153 DNA see 1d Accession lence: 149-' 11 AAAGCGCGGA GAGTGGAAG GAGTGGAAAG CGCAGCCCG GAGATGAAAG CGAAGTGATT TGCACTGTAC GGTAGCGTCT AATAATTATA GACATTACAA GACTATGGAG GACTATGGAG	LKUVKPFGDS VKWHTAVTYV FFTCFLPYHL FFRRLFKKSN Quence 1 #: D80008 739 21) GCGGAGGCCG GAGCCCAGAT CGTCGCAT TGATACCAAC CATACCTGTA TGATACCAAC CATACCTGTA TGCAAATGC AAGATCTCA AGGATATGAA AGATTGTAA	RMYSITFTKV NSCLFVAVLV CRIPFTFSHL IRTRSESIRS 1.1 31 AGGCGAGAGC ACCATTTTGG GTTCTGCGAA ACCAGCTTC ACAAAACCAG TATCAAATTT TGACCGCTTG ATTACGATTT TGCTACTATA ACCACCAAAA ACCACCAAAA ACCACCAAAA ACCACC	LSYCWYIMA ILIGCYIAIS DRLLDESAQK LQSYRRSEVR 41 CTGGCGCTGT CGTGAGAGCT AAAGCCATGG TCTGATGTGA TCTTGATGTGA CCACACTGTT ATGAGGTCA CACATGGCT ATGAGGTCA AGCCTATATA ACTTCAGTCC	VISLPNIILT RYIHKSSRQF ILYYCKETTL IYYDYTDV 51 AGGACTAGAA GGTGGTTGGC AACTGATCGG GACTCAGACA ATGAAGCAAA CTCTGTTAAG GAGCACTCAG CTGAAGAAAT TGGGAGGAGA TTGAAGTCCG TTATAAAAAA	60 120 180 240 300 420 480 540 660
50 55 60	FYANMYTSIV NGQPTEDNIH ISGSSRKRKH PLSACNVCLD Seq ID NO: Nucleic Ac: Coding sequ 1 GTTCGGCGCC CGAAAGGAGT AAGGCCGCGG CGAACTGCAT AGTTCTGGAG GTCAGGGGAATAT GGAGTAGTTT TGAAGGTTTT GGAAGGTTGT GTGTCTAAAA AAATAGCCAG	FLGLISIDRY DCSKLKSPLG NQSIRVVVAV PILYFFMCRS 153 DNA see id Accession lence: 149- 11 AAAGCGCGGA GAGTGGAAG GCGCGCCG GAGATGAAAG CGAAGTGAAAG GGTAGCGTCT TGCACTGTAG GGTAGCGTCT AATAATTATA GACATTACAC GACTATGGAG GACATTACAC CACTTATGGA	LKVVKPFGDS VKWHTAVTYV FFTCFLPYHL FFRRLFKKSN Quence 1 #: D80008 739 21 GCGGAGGCCG GAGCCCAGAT CGTCCGCCAT AAGGGCAACT CTTTGTATGA TGATACCTGTA TGCAAATGC AAAGATCTCT AGGATATGAA AATTTGAAGT CTCGATGGAA ACTTGAAGT	RMYSITFTKV NSCLFVAVLV CRIPFTFSHL IRTRSESIRS 1.1 31 AGGCGAGAGC ACCATTTGG GTTCTGCGAA GCCTGCCTTC ACAAAACCAG TATCAAATTT TGACGCTTG ATTACGATTT TGCTACTTAT ACCACCAAAA TGATGATGAGCAG ATGTGAGCAG	A1 CTGGGGCTGT CGTGAGAGCT AAAGCCATGG CCTCGGAGAGCT AAAGCCATGT CTTCGGATCA CACATGGCTG ATGAGGTCA AGCCTATATA ACTTCAGTCC CTGATCACAC CTGATCACAC CTGATCACAC CTGATCACAC CTGATCACAC CTGATCACAC CTGATCACAC CTGATCACAC CTGATCACAC CTGATCACAC CTGATCACAC CTGATCACAC CTGATCACAC	VISLPNIILT RYIHKSSROF ILYYCKETTL IYYDYTDV 51 AGGACTAGAA GGTGGTTGGC AACTGATCCG GACTCAGACA ATGAAGCAAA CTCTGTTAAG GAGCACTCAG CTGAGAGAAT TGGGAGGAGA TTGAAGTCCT TTGTAAGAGCAAA AAGGAGTCCT	180 240 300 120 180 240 360 420 480 600 600 720
50 55 60	FYANMYTSIV NGQPTEDNIH ISQSSRKRKH PLSACNVCLD Seq ID NO: Nucleic Ac: Coding sequ [] GTTCGGCGCC CGAAAGGAGT AAGGCCGCG GCAACGGAGT AGGTCTGGAG GTCAGGTGGA AAATCGAACC ATGGGAATAT GGAGTGGTTT GTGTCTAAAA AAATAGCAG GGAGGAGTC	FLGLISIDRY DCSKLKSPLG NQSIRVVVAV PIIYFFMCRS 153 DNA see id Accession lence: 149-' 11 AAAGCGCGGA GAGGGCCGA GAGTGGGAAG CGGGCCCG GAGATGAAAC CGAAGTGATT TGCACTGTAG GGTAGCGTCT AATAATTATA GACATTACAC GACTATGGAG CACTTTTTAC CCTGTGAGAG CACTTTTTAC	LKUVKPFGDS VKWHTAVTYV FFTCFLPYHL FFRRLFKKSN Quence 1 #: D80008 739 21 GCGGAGGCCG GAGCCCAGAT CGTCCGCCAT AAGGGCAAC CTTTGTATGA TGATACCAAC CATACCTGTA TGCCAATGC AAAGATCTCT AAGGATTGTATGAA AATTTGAAGT ACTGGATGGAA AATTTGAAGT CTCGATGGAA AATTTGAAGT CATGCCGA	RMYSITFTKV NSCLFVAVLV CRIPFTFSHL IRTRSESIRS 31 31 31 31 CCATTTGG GTTCTGCGAA GCCTGCCTTC ACAAAACCAG TATCAAATTT TGACCGCTTG ATTACGATTT TGACTACTTAT ACCACCAAAA TGATGATGGC ATGTGAGCAG GGCACTTCCA	LSYCWYIMA ILIGCYIAIS DRLLDESAQK LQSYRRSEVR 41 CTGGCGCTGT CGTGAGAGCT AAAGCCATGG AACGAGGATG TCTGATGTGT CTTCGGATCA CACATGGTA ACATGGCTG ATGAGGTCA AGCCTATATA ACTTCAGTCC CTGATCAGC CGGCTTCACTC	VLSLPNIILT RYIHKSSROF ILYYCKEITL IYYDYTDV 51 AGGACTAGAA GGTGGTTGGC AACTGATCCG GACTCAGACA ATGAAGCAAA CTCTGTTAAG CTGAAGAAA TCGGAGGAGA TTGGAGGAGT TTGAAGTCCG TATTAAAAA AAGGAGTCCT TATTAAAAAA AAGGAGTCCT TATCATGGA	60 120 180 240 300 420 480 540 660 720 780
50556065	FYANMYTSIV NGQPTEDNIH ISGSSRKRKH PLSACNVCLD Seq ID NO: Nucleic Ac: Coding sequ 1 GTTCGGCGCC CGAAAGGAGT AAGGCCGCGG GTCAGGTGGA GTCAGGTGGA GTCAGGTGGA AAATCGACGC ATGGGAATAT GGAGTTGTT TGAAGGTTT TGAAGGTTTT TGAAGGTTTT GGAGGTAAA AAATAGCCAG GGAGCACATC CTCCTCTGTT TAGACCATTGT TTAGACCATTGT TTAGACCATTGT	FLGLISIDRY DCSKLKSPLG NQSIRVVVAV PIIYFFMCRS 153 DNA see id Accession ence: 149-' 11 AAAGCGCGGA GAGGGCCGA GAGTGGAAAG CGAGTGGAAG CGAGTGATAT GGACATTATA GACATTATA GACATTATA GACATTATA CACTTATGA CACTTTTTAC CACTTTTTAC CACTCTCT TTAAGATAAAA	LKUVKPFGDS VKWHTAVTYV FFTCFLPYHL FFRRLFKKSN Quence #: D80008 739 21 GCGGAGGCCG GAGCCCAGAT CGTCCGCAT AAGGGCAACT CTTTGTATGA TGATACCTGTA TGCAAATGC AAGATCT AGGATATGAA AATTTGAAGT CTCGATGGAA CTTGGATGGAA CATGCGCCGA CATGCACCCC TAAGAATACC TAAGAATACC TAAGATACCTACC TAAGAATACC TAAGAATACC TAAGAATACC TAAGAATACC TAAGAATACC	RMYSITFTKV NSCLFVAVLV CRIPFTFSHL IRTRSESIRS .1 31 AGGCGAGAGC ACCATTTGG GTTCTGCGAA GCCTGCCTTC ACAAAACCAG TATCAAATTT TGACCGCTTG ATTACATTAT ACCACCAAAA TGATGATTAT ACCACCAAAA TGATGATGAGCA CTTCACTTCC CTTCACCTCC TGGCTAAGAA	A1 DRILDESAQK LQSVRRSEVR A1 CTGGCGCTGT CGTGAGAGCT AAAGCCATGGT CTTCGGATCA CACATGGTG ATGAGGTCA AGCCTATATA ACTTCAGTCC CTGATCAGC CTGATCAGCC CTGATCAGC CCTCTTCACTC CTGATCATT GTATAATTTG	VISLPNIILT RYIHKSSROF ILYYCKETTL IYYDYTDV 51 AGGACTAGAA GCTGGTTGGC AACTGATCCG GACTCAGACA ATGAAGCAAA CTCTGTTAAG GAGCACTCAG CTGAAGAAAT TGGAGGAGA TTGAAGTCCT AACTCATGAA TATAAAAAA AAGGAGTCCT AACTCATGAA CTTATGAAGCTA CTAACTATTA	180 240 300 120 180 240 360 420 540 660 720 780 840 900
50 55 60	FYANMYTSIV NGQPTEDNIH ISGSSRKRKH PLSACNVCLD Seq ID NO: Nucleic Ac: Coding sequ 1 GTTCGGCGCC CGAAAGGAGT AAGGCCGCGG CGAGCTGCAT AGTTCTGGAG GTCAGGTGGAA AAATCGACGC ATGGAATAT TGAAGGTTTT TGAAGGTTTT TGAAGGTTTT GGGGTTAAAA AAATAGCCAG GGAGCACATC CTCCTCTGTA TAGACATTGT TAGACATTGT AGGACTTTCT AGGACTTTCT TGAGCATTTTT TGAGACATTGT AGGACTTTCTT	FLGLISIDRY DCSKLKSPLG NQSIRVVVAV PILYFFMCRS 153 DNA see id Accession lence: 149- 11 AAAGCGCGGA GAGTGGGAAG CGCGCGCG GAGATGAAG CGAGTGGTT TGCACTGTAG GACTATGAG GACTATGAGA CACTTTTAC CACTTTTAC CACTTTTAC CACTTCTTA	LKVVKPFGDS VKWHTAVTYV FFTCFLPYHL FFRRLFKKSN Quence 1 #: D80008 739 21 GCGGAGGCCG GAGCCCAGAT CGTCCGCCAT AAGAGCCAACTCC TAAGAATACTAA CATGCAACTCT AGGATATGAA TGCCAAATGC AAGATATGAA CATGCACCCCACCC	RMYSITFTKV NSCLFVAVLV CRIPFTSHL IRTRSESIRS 1.1 31 AGGCGAGAGC GTTCTGCGAA GCCTGCCTTC ACAAAACCAG TATCAAATT TGACGCTTG ATTACGATT TGATACTATT ACCACCAAAA TGATGATGG GGCACTTCCA CTTCACCTCC TGGCTAAGAA TTCTTCCTAC	A1 CTGGCGCTGT CGTGAGAGCT AAAGCCATGG ACAGGGATGT CTTCGGATCA ACTGTCATCAGGCC ATCAGGCC ATCAGGCC ATCAGGCC ATCAGGCC ATCAGGCC ACATGCT ATCAGGCC ACCTGTT CTTCGGATCA ACTTCAGTCC CTGATCAGC GGCTTCACTC CTGATCAGC GGCTTCACTC CTCTTTGATT GTATAATTTG	VISLPNIILT RYIHKSSROF ILYYCKETTL IYYDYTDV 51 AGGACTAGAA GGTGGTTGGC AACTGATCCG GACTCAGACA ATGAAGCAAA CTCTGTTAAG GAGCACTCAG CTGAAGAAAT TGGAAGCACT TAGAAGCACT AACTCATGA TTAGAAGCT CTAACTATTA TTTTGGTTTT	180 240 300 60 120 180 240 360 420 480 540 600 660 720 780 840 960
50556065	FYANMYTSIV NGQPTEDNIH ISGSSRKRKH PLSACNVCLD Seq ID NO: Nucleic Ac: Coding sequ 1 GTTCGGCGCC CGAAAGGAGT AAGGCCGCGG GTCAGGTGGA AGATCGAGA GTCAGGTGGA GTCAGGTGGA AGATCGAGT GGAGTGGTT TGAAGGTTT TGAAGGTTT TGAAGGTTCT GGACTCCTCTGTA AAATAGCCAG GGACACATC CTCCTCTGTT AGGACTTTCT GGTTTTTGTAGA AGTCTCTCCCA	FLGLISIDRY DCSKLKSPLG NQSIRVVVAV PIIYFFMCRS 153 DNA see id Accession ence: 149-' 11 AAAGCGCGGA GAGTGGAAG GAGGGCCCG GAGATGAAAG CGAAGTGATT TGCACTGTAG GACTATCAG CACTTTTAC CACTTTTAC CACTCTCT TTAAGATAAC TTTTTTAATG GACTGTCTCA TTTTTAATG GACTGTCTCA CCCTCACTCT TTAAGATAAC CTCTACTCTCT TTAAGATAAC CTCTACTCTCT CACTGTCT T CACTGTCT ACTGTC CACTC CACTGT CACTGT CAC	LKUVKPFGDS VKWHTAVTYV FFTCFLPYHL FFRRLFKKSN Quence #: D80006 739 21 GCGGAGGCCG GAGCCCAGAT CGTCGCCAT AAGGGCAACT CTTTGTATGA TGATACCTATA TGCAAATGC AAGATCT AGGATATGAA AATTTGAAGT CTCGATGGAA CATGCGCCGA CATGCGCCGA TTGATGAA CATGCACCACT TTGACACTA TTGTACACTA CTATGTTGCC TAAGAATACT TTGTACACTA CTATGTTGCC TCAAGATGTT TCTACACTA CTATGTTTGCC TCAAGATGTT TCTAAGTTTT TCTAAGTTT TCTAAGTTTT TCTAAGTTTT TCTAAGTTTT TCTAAGTTTT TCTAAGTTTT TCTAAGTTTT TCTAAGTTTT TCTAAGTTT TCTAAGTTTT TCTAAGTTTT TCTAAGTTTT TCTAAGTTTT TCTAAGTTT TCTAAGTGTT TCTAAGTGTT TCTAAGTGTT TCTAAGTGTT TCTAAGTGTT TCTAAGTGTT TCTAAGTGTT TCTAAGTGTT TCTAAGTGTT TCTAAGTGTT TCTAAGTGTT TCTAAGTGTT TCTAAGTGTT TCTAAGTGTT TCTAAGTGTT TCTAAGTGTT TCTAAGTGTT TCTAAGTGTT TCTAAGTGTT TCTAAGTT TCTAAGT TCTAAGTT TCT	RMYSITFTKV NSCLFVAVLV CRIPFTFSHL IRTRSESIRS 1.1 31 AGGCGAGAGGC ACCATTTGG GATTCTGCGAA GCCTGCCTTC ACAAAAACCAG TATCAAATT TGACCGCTTG ATTACGATTAT ACCACCAAAA TGATGATTCTACCTCC CTTCACCTCC CTTCACCTCC CTGCCTAAGAA TTCTTCCTAC CAAGCTGGTC CAAGCTGGTC GAGATCACAA	A1 DRILDESAQK LQSVRRSEVR A1 CTEGGCGCTGT CGTGAGAGCT AAAGCCATGGT CTTCGGATCA CACATGGTC ATGAGGTCA AGCCTATATA ACTTCAGTCC CTGATCAGC CTGATCAGCT CTTTTGTT GTATATTTG GTTATATTTT GTCTTTTTTTGG TCAAACTCCT GCGTAGACCA	VISLPNIILT RYIHKSSROF ILYYCKETTL IYYDYTDV 51 AGGACTAGAA GGTGGTTGGC AACTGATCCG GACTCAGACA ATGAAGCAAA CTCTGTTAAG GAGCACTCAG CTGAAGAAAT TGGAGGAGA TTGAAGTCCT AACTCATGAA TTATAAAAAA AAGGAGTCCT AACTCATGAA CTTATTATGATTTT TTTTGGTTTT GGCCTCAAGC CTGCACCCGG	180 240 300 60 120 180 240 360 420 540 660 720 780 840 960 960 1020
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5055606570	FYANMYTSIV NGQPTEDNIH ISGSSRKRKH PLSACNVCLD Seq ID NO: Nucleic Ac: Coding sequ 1 GTTCGGCGCC CGAAAGGAGT AAGGCTGCAT AAGTTCTGGAG GTCAGGTGGA AAATCGACGC ATGGGAATAT GGAGTGGTT TGAAGGTTT TGAAGGTTT TGAAGGTTT TGAAGGTTT TGAAGGTTT TGATTTTT AGACATTGT AGGACTTCCT GTTTTTTTTAGA AGTCCTCCCA CCCCTACTCC GTGTGTTTTTT TTGGCTGGAC	FLGLISIDRY DCSKLKSPLG NQSIRVVVAV PIIYFFMCRS 153 DNA see id Accession ence: 149- 11 AAAGCGCGGA GAGTGGAAG GAGTGGAAG GCGCGCCG GAGATGAAAG CGACTGTAT GGCATTTAC GACTATTAC GACTATTAC GACTATGAG CACTTTTAC CTGTCATGAC CTCACTCTC TTAAGATAAC TTTTTTAATG GACTGTCTC TTAGCTTC TTAGCTTC TTATGCTTC TTTTTCTAAT TAAATGAAAG AGGAAGAAGG	LKUVKPFGDS VKWHTAVTYV FFTCFLPYHL FFRRLFKKSN Quence #: D80008 739 21 GCGGAGGCCG GAGCCCAGAT CGTCGGCAT AAGGGCAACT CTTTGTATGA TGATACCTATA TGCCAAATGC AAGATCTCT AGGATATGAA AATTTGAAGT CTCGATGGAC CTTGGACGAA CTTGGACGAA CTTGGACGAA CTTGGACGAA CTTGGATGAGA CATGCGCGA CATGCGCCGA CCACACTCC TAAGAATACT TTGTACACTA CTATGTTGCC TCAAAGTGTT AAGCTGTATC TAAACATGGT TAAACATCGTT TAAACATCGTT TAAGCTGTT	RMYSITFTKV NSCLFVAVLV CRIPFTSHL IRTRSESIRS 1.1 31 AGGCGAGAGC ACCATTITGG GTTCTGCGAA GCCTGCCTTC ACAAAACCAG TATCAAATTT TGACCGCTTG ATTACGATTAT ACCACCAAAA TCATCACATTCA CACACCAAAA TCATCACCTCC TGGCTAAGAA TTCTTCTCCTAC CAAGCTGGTC GAGATCACAG TGACTACACA TGACTAGAC TGACTAGACA TTCTTCTCTAC CAAGCTGGTC GAGATCACAG TGTAATCACA TGCATTTGAA TGATTGAA TGCTTGTTT	A1 CTGGCGCTGT CTGGAGAGCT AAAGCCATGGT CTTGGATCA CACATGGTCA ATGAGTCA ACTCAGTCA CTGATCAGTCA CTGATCAGTCA CTGATCAGTCA CTGATCAGTCA CTCATTCAGTCA CTCATTCAGTCA CTCATTCAGTCA CTCTTTATTTGG TCAACTCCT GTATAATTTG TCTTTTTTTGG TCAACTCCT GCGTGAGCCA GCATTCCTAA TCTCTTAAAT TTCTTGGTCA	VISLPNIILT RYIHKSSROF ILYYCKETTL IYYDYTDV 51 AGGACTAGAA GGTGGTTGGC AACTGATCCG GACTCAGACA ATGAAGCAAA CTCTGTTAAG GAGCACTCAG CTGAAGAGAAA TTGAAGTCCT AACTCATGAA TTAGAAGTCCT AACTCATGAT TTTGGTTTT GGCCTCAAGC CTGCACCCGG AGTTGTTACA AAGCAGCCCCGG AGTGTTTCAC GTGTATTGTAT	180 240 300 60 120 180 240 300 360 420 720 780 960 1020 1080 1140 1260
50556065	FYANMYTSIV NGQPTEDNIH ISGSSRKRKH PLSACNVCLD Seq ID NO: Nucleic Ac: Coding sequ 1 GTTCGGCGCC CGAAAGGAGT AAGGCCGCGG CGAACTGCAT AGTTCTGGAG GTTCAGGAG ATGGAATAT GGAGTTGTT TGAAGGTTTT TGAAGGTTTT TGAAGGTTTT TGAACATTGT AGGACTTCTT ATGACATTGT AGGACTTCTT GTTTTTTAGA AGTCCTCCC GTGTGTTTTT TTGGCTGAA AGTCCTCCC CCCTACTCC CCCTACTCC CAAGTTGAAC CCAAGTTGAAC CCCCTACTCC CTGTGTTTTT TTGGCTGAAC CAAGTTGAAC CAAGTTGAAC	FLGLISIDRY DCSKLKSPLG NQSIRVVVAV PILYFFMCRS 153 DNA see id Accession lence: 149-' 11 AAAGCGCGGA GAGTGGGAAG GAGTGGAAG GAGATGAAAG GGAGTGGAAG GGTAGCGTCT AATAATTATA GACATTACAC CACTATGGA CACTTTTTAC CACTCTCT TTAAGATAAC CTCACTCTCT TTAAGATAAC CTTACTCTCT TTAAGATAAC CTTACTCTCT TTAAGATAAC CTTAGCTTC CTTTTTTAAT GACATGACAAG ACGAAGAAGAAG AGGAAGAAGAAG AGGAAGAAGT	LKUVKPFGDS VKWHTAVTYV FFTCFLPYHL FFRRLFKKSN TUENCE 1 #: D80008 739 21] GCGGAGGCCG GAGCCCAGAT CGTCCGCCAT AAGGGCAACT CTTTGTATGA TGCAAATGC AAGATACTA AGTTTGAAGT CTCAGATACTA CTAGATACTA TTGTACACTA CTAAGATACT TAGATACTA TTGTACACTA TTGTACACTA CTAAGTTTTCTACACTA TAGATACTA TTGTACACTA TTGTACACTA TTGTACACTA TTGTACACTA TTGTACACTA TTGTACACTA TTGTACACTA TTAGATGCT TAAACATGCT TAAACATGCT TAAACATGCT TAAACATGCT TTGTACACTA TTGTACACTA TTGTACTTATCT TAAACATGCT TAAACATGCT TTGTACACTA TTGTACACT	RMYSITFTKV NSCLFVAVLV CRIPFTSHL IRTRSESIRS 1.1 31 AGGCGAGAGC ACCATTTGG GTTCTGCGAA GCCTGCCTTC ACAAAACCAG TATCAAATTT TGATACTTAT ACCACCAAAA TGATGATGAG ATTGAGCAG TATCAAGATTAT ACCACCAAAA TGATGATGAG TATCAACTCCA CTTCACCTCC CTTCACCTCC CTTCACCTCC CAGGATAGAA TTCTTCCTAC CAAGCTGGTC GAGATCACAG TTCTTCTAC CAAGCTGGTC CAGGTTAGAA TTCTTCTAC CAAGCTGTTC CAGGATCACAG TGTAATCACA TGATTTGAA GTGTTCTTGTT CATTTTCAAA	A1 CTGGCGCTGT CGTGAGAGCT AAAGCCATGT CTTCGGATCA ACCTTCAGTCA ACCTTCAGTCA CTGATCAC ACCTTTCAGTCA CTGATCAC CTGATCAC CTCATCAC CTGATCAC CTCATTCAGTCA CTGATCAC CTCATTCAGTCA CTGATCACC CTCATTCAGTCA CTCATTCAGTCA CTCATTCAGTCA CTCATTTAATT CTTTTTTTTG CTATAATTG TCTTTATTT TCTTTTATTT TCTTTTATTT TCTTTTATTT TCTTTTATTT TCTTTTAATT TCTCTTAAAT TTCTGGTCAT TTCACATGCAT	VISLPNIILT RYIHKSSROF ILYYCKETTL IYYDYTDV 51 AGGACTAGAA GGTGGTTGGC AACTGATCCG GACTCAGACA ATGAAGCATAA CTCTGTTAAG GAGCACTCAG ATGAAGTCCG TATTAAAAAA AAGGAGTCCT AACTCATGGA TTAGAAGCTT TTAGAAGCTT TTAGAAGCTT TTAGAAGCTA TTAGAAGCTA CTAACTATTA TTTTGGTTTT GCCCTCAACCGG AGTTGTTACA AAGCAGTCAC GTGTATTGTA GTGAAGGTTAC	180 240 300 120 180 240 360 420 480 540 660 660 720 780 840 900 960 1020 1020 1140 1200 1200 1320
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505560657075	FYANMYTSIV NGQPTEDNIH ISQSSRKRKH PLSACNVCLD Seq ID NO: Nucleic Ac: Coding sequ (Coding FLGLISIDRY DCSKLKSPLG NQSIRVVVAV PIIYFFMCRS 153 DNA see id Accession lence: 149-' 11 AAAGCGCGGA GAGTGGAAG GAGTGGAAAG GCGCGCCG GAGATGAAAG CCAATTACA GACTATTACA GACTATTACA CACTATTACA CACTATTACA CACTATTACA CACTATTACA CACTATTAATA GACATTATAA GACATTATAA GACATTATAA GACATTATAA GACATTATAA GACATTATAA GACATTATAA GACATTATAA GACATTATAA GACATTATAA GACATTATAA GACATTATAA GACATTATAA GACATTATAA GACATTATAA CACTTTTAAT TAAATGAAAG AGCAGAATTT AAAATTATAA GGAAGAAGA GGGAGGACAC TGGGGTGATC AACTTTGTAC TGGGGTGATC AACTTTGTAC CGTCTTTAAT	LKUVKPFGDS VKWHTAVTYV FFTCFLPYHL FFTCFLPYHL FFRRLFKKSN Quence 1 #: D80008 739 21) GCGGAGGCCG GAGCCCAGAT CGTCGGCAT TAGATACCAAC CATACCTGTA TAGCAATGC AAGATATCAA CATGACTGAA CATGACTGA CATGACTGA CATGACTGA CATGACTGA CATGACTGA CATGACTGA CATGACTGA CATGACTGA CATGACTGA CATGACTGA CATGACTGA CATGACTGA CATGACTGA CATGACTGA CATGACTGA CATGACTGA CATAGATACA AATTTGACACT TAGATCCTGT TAGATCCTGT TAGATCTGT CTGAGATACA AATTTTTCCT ACCAGTATCA AGTTTTTCCT GGTATTTTCCTT GGTATTTTCCTT GGTATTTTCCTT GGTATTTTCCTT GGTATTTTCCTT GGTATTTTCCTT GGTATTTTCCTT GGTATTTTCCTT GGTATTTTCCTT GGTATTTTTCCTT GGTATTTTTCCTT GGTATTTTTCCTT GGTATTTTTCTT TAGATCTTT TCTTT TAGATTTTTCTT TAGATTTT TAGATTTT TAGATTTT TAGATTTT TAGATTT TAGATTT TAGATTT TAGATTT TAGATTT TAGATT	RMYSITFTKV NSCLFVAVLV CRIPFTSHL IRTRSESIRS 1.1 31 AGGCGAGAGC ACCATTITGG GTTCTGCGAA ACCAGCTTC ACAAAACCAG TATCAAATTT TGTACTATA ACCACCAAAA TCATGAGCAC ATGAGCACAAAA CTTCACCTCC TGGCTAAGAA TTCTTCTCCTAC CAAGCTGGTC GAGATCACAG TATATCACATTC GAGATCACAG TATTATTGAA TTTTTCACAT CCAACTTTGAA GTTTAATGAC TGCACATTTC CAACTTTGAA GTTTAATGAC GTGCACATTTC CACTTTGAA ACTTTTAATGAC GAGATTCAGA AAACACTTAAGA AAACACTTAAGAA AAACACTTAAGAA AAACACTTAAGAA AAACACTTAAGAA AAACACTTAAGAA AAACACTTAAGAA AAACACTTAAGAA AAACACTTAAGAA AAACACTTAAGAA AAACACTTAAGAA AAACACTTAAGAA AAACACTTAAGAA AAACACTTAAGAA AAACACTTAAGAA AAACACTTAAGAA AAACACTTAAGAA AAACACTGAA	LSYCWYIMA ILIGCYIAIS DRILDESAQK LQSYRRSEVR A1 CTGGCGCTGT CGTGAGAGCT AAAGCCATGGT TCTGATGTGA TCTGATGTGA ACTACAGCTG ATGAGGTCA ACTCAGTCC CTGATCAGCC TGATCAGCC GCCTTCACTT GTATATATTG TCTTTTTTGG TCAAACTCCT GCGTGAGCCA GCATTCCACT GCGTGAGCCA GCATTCCTAC TCACATGCAT TCACATGCAA ATACTAAATTT CACCATGCTA ATACTAATTT CACCATGCTA ACTCATGCAA ACTCATGCAAA ACACTAAATTT CACCATGGTG AGGGGACAGT TTGACTGAAA CATTTTAAAT	VISLENTILT RYIHKSSROF ILYYCKETTL IYYDYTDV 51 AGGACTAGAA GGTGGTTGGC AACTGATCCG GACTCAGACA ATGAAGCAAA CTCTGTTAAG CTGGAGGAAA TTGAGGTCCG TATTAAAAAA AAGGAGTCCT AACTCATGGA ATCATGGA ACTCATGGA CTGACCAAGC CTGACCAAGC CTGACCAGG AGTTGTTACA AACTATTA TTTTGGTTTT GGCCTCAAGC CTGCACCCGG AGTTGTTACA AAGCAGTCAC GTGATCTGA GCCTGATCCGG AGTTGTTACA ATCATCTGG GCACTCGG GAAATTGGGG CTGGTGTGG GAAATTGGGG AGTCACATGA TTTGATGAAAA	180 240 300 60 120 180 240 360 420 780 960 960 1080 1140 1260 1320 1380 1440 1560 1620	
505560657075	FYANMYTSIV NGQPTEDNIH ISQSSRKRKH PLSACNVCLD Seq ID NO: Nucleic Ac: Coding sequence (GTTCGGCGCC CGAAAGGAGT AAGGCCCGG GCAAGGAGT AAGTCTGGAG GTTCTGGAG GTTCTGAGTGAA AAATCGAAGTTT TGAAGGTTTC TGGAGGTTTC TGGACATCC CTCCTCTGTA TAGACATTCT GTTTTGTAGA AGTCCTCCCC CCCTACTCC CCCTACTCC CCCTACTCC CAGTTTTT TTGGCAA TATTTGGGAA CTTTTGGGAA CTTTTGGGAA CTTTTTGGCAA CTCAGTTGAT TCCCAGTTTAT TCCCAGTTAT TCCCAGTTTAT TCCCAG	FLGLISIDRY DCSKLKSPLG NQSIRVVVAV PIIYFFMCRS 153 DNA see id Accession ence: 149- 11 AAAGCGCGGA GAGTGGAAG GAGGCGCCG GAGATGAAAG CGCAGCCCG GAGATGATAT TGCACTGTAG GGTAGCGTCT AATAATTATA GACATTACAC CACTTTTTAC CTTCATGAG CACTTTTTAAT GACTGCTCT TTAAGATAAC CTTTATGTAC CTTAGCTCC TTTTTCTAAT GACTGATAG AGCAAGAGAG AGCAAGAGAG AGCAAGAGAC CTTTTTCTAAT CACTTTTTAAT GGAAGGACAC TGGGGTGATC TAAATGAAAG AGCAAGAGAC CTGGGTGATC CTTTTTCTAAT CGGAAGGACAC TGGGGTGATC CACTTTTTAAT CGTCTTTTAAT CGTCTTTTAAT CGTCTTTTAAT CGTCTTTTAAT CCATTTTTTT	LKUVKPFGDS VKWHTAVTYV FFTCFLPYHL FFRRLFKKSN Quence #: D80008 739 21 GCGGAGGCCG GAGCCCAGAT CGTCGCCAT AAGGGCACT CTTTGTATGA TGATACCAAC CATACCTGTA TGCAAATGC AAAGATCTCT TAGATAGAA AATTTGAAGT CTCGATGGAA AATTTGAAGT CTCAATGGAATAC TGAAGATACT TTAGACTAT TGAAGATACT AAGCTGTAT TAGAATACT AAGCTGTAT AAGCTGTAT AAGCTGTAT AAGCTGTTT AAGCTGTTT AAGCTGTTT AAGCTGTTT AAGCTGTTT TAGAATACA ACTTTTCCCT GGTATGTTT TTTTTTTTTT	RMYSITFTKV NSCLFVAVLV CRIPFTFSHL IRTRSESIRS 31 31 AGGCGAGAGC ACCATTTGG GTTCTGCGAA ACCAGTTTTGACGCTTG ATTACAATTT TGACCGCTTG ATTACGATTT TGACTACTATA ACCACCAAAA TGATGATGGC ATGTGAGCAG GGCACTTCCA CCTCCACTCC TGGCTAGCAA TTCTTCCTAC CCAAGCTGGT CAAGCTGGT CAAGCTGGT CAAGCTGGT CAAGTTTGAA GTTAATCACA TTCTTCTAC CCACTTTGAA GTTTAATGAC GTCACTTTGAA GTTTTAATGAC TGCACATTTGA AACAGCTGA AAACAGCTGA TGGGTGTTGAA CAGTTTGAA CAGTTTAGAA CTTTTAGAAG CTTTCTAGAAG CTTTTAGAAG CTTTTTAGAAG CTTTTAGAAG CTTTTTAGAAG CTTTTAGAAG CTTTTTAGAAG	LSYCWYIMA ILIGCYIAIS IRILDESAQK IRICORYRSEVR AI CTGGCGCTGT CGTGAGAGCT CGTGAGAGCT TCTGGGTCATGG AACGAGGATG TCTGATGTGA ACCATGGTA ACTTCAGTCC CTGATCAGC CACTGTATATA ACTTCAGTCC CTCTTTGATT GTATCAGTCC CTCTTTGATT GTATAATTTG TCTAACTCCT GCGTGACCAG CATTCCTAC TCTCTTAAT TCCTGGATCAC CACATGCTG TCAACTCCTAC TCAACTCCTAC TCAACTCCTAC TCAATGCAA ATACTAATTT CACCATGGTG AGGGGACAGT TTCACTGAAA CATTTTAAAT ATCCGAGAAA ATCTTAAAT ATCCGAGAAA CATTTTAAAT ATCTGATAATT	VISLENTILT RYIHKSSROF ILYYCKEITL IYYDYTDV 51 AGGACTAGAA GGTGGTTGGGC AACTGATCCG GACTCAGACA ATGAGCAAA TCGAGCACA ATGAGCACA ATGAGCCCG TATTAAAAAA AAGGAGTCCT TATGAGAGTCT TTTGGTTTT GGCCTCAAGC CTGCACCCGG AGTTGTACA AAGCAGTCAC GTGTATTGTA GTGAAGATGA ATCATCTGGC GGTGATGTACA ATCATCTGGC GCTGTGTACC GTGTATTGTA TTGTATTGTA	180 240 300 60 120 180 240 420 480 600 660 6720 780 840 960 1020 1140 1200 1140 1320 1380 1440 1560 1620 1620 1620 1740
50 55 60 65 70 75	FYANMYTSIV NGQPTEDNIH ISQSSRKRKH PLSACNVCLD Seq ID NO: Nucleic Ac: Coding sequ (Coding FLGLISIDRY DCSKLKSPLG NQSIRVVVAV PIIYFFMCRS 153 DNA see id Accession lence: 149- 11 AAAGCGCGGA GAGTGGAAG GAGGGCCGG GAGATGAAAG CGCAGCCCG GAGATGAAAG CGACTATTAC GACTATTAC GACTATTAC CTGTCATTAGAC CTCACTCTC TTAAGATAAC CTCACTCTC TTAAGATAAC CTCACTCTCT TTAAGATAAC CTCACTCTCT TTAAGATAAC CTCACTCTCT TTAAGATAAC CTCTACTCTCT TTAAGATAAC CTCTACTCTCAT TAAATTATAA GACAATTTTT AAATTATAA GACAAGAAGA GGCTGAATTT AAATTTTTAA CGGTTGATT AAATTTTTT ACGTTTGTAC CTCTTTTTTAAT TCGTTTGTAC CTCTTTTTTTAAT TCGTTTGTTC ACAATTTTTT TATGACCGT	LKUVKPFGDS VKWHTAVTYV FFTCFLPYHL FFTCFLPYHL FFRRLFKKSN Quence 1 #: D80008 739 21) GCGGAGGCCG GAGCCCAGAT CGTCGCAGAT CGTTCGCATGA TGATACCAAC CATACCTGTA TGCAAATGC AAGATTCTA AGGATATGA AATTTGAAGT CTCATGAGAT CTAGATGCCCGA CCACCCC TAAGAATACT TTGTACACTG TAGATACTG TAGATCTGT TAGATCTGT TAGATCTGT TAGATCTGT TAGATCTGT TAGATCTGT TAGATCTGT TAGATCTGT TAGATCTGT TAGATCTGT TTGAGATTA ACATGGATTT ACCAGTATCA AGTTTTTCCT TGTTTTTTT TTTTTTTTTT	RMYSITFTKV NSCLFVAVLV CRIPFTFSHL IRTRSESIRS 1.1 31 AGGCGAGAGC ACCATTTGG GTTCTGCGAA ACCAGCTT TGACCGCTT ACCACCACAA ACCATTTTGG GTTCTGCTAC ACTTCACTCC TGCTAGCAC ACTTCACCTCC TGCTAGCAC TTCTCCTAC CAAGCTGGTC GAGATCACA GTTTATCACA GTTTATTGAC GTTTATTCACA GTTTATTCACA GTTTATTCACA GTTTATTCACA GTTTATTCACA GTTTATTCACA GTTTATTCACA GTTTTATTCACA GTTTTCACA GTTTTATTCACA GTTTTCACA GTTTTCACA GTGCTTTGGA AAACAGCTGA TGCGTGTTGC TTCTAGGA TGGGTGTTGC TTTTTTTTTT	LSYCWYIMA ILIGCYIAIS DRILDESAQK LQSYRRSEVR A1 CTGGCGCTGT CGTGAGAGCT CGTGAGAGCT CGTGAGGAGCT TCTGATGTGA ACACTGTT CTTCGGTCC CTGATCAGTCC CTGATCAGTCC CTGATCAGTCC CTCTTTTATT GTATAATTTG TCAACTCCT GCGTGAGCCA GCATTCCTAC TCACTGATC TCACATGCTA ATTCTGGTCAT TCACATGCAA ATACTAATTT CACATGCAA ATACTAATTT CACCATGGTG AGGGGACAGT TTGACTGAAA TCTTAAATT TCCGAGAAA ATACTAAATT CACCAGGAAA ATGTTATAAATT GTTTTTTTTTT	VISLENTILIT RYIHKSSROF ILYYCKETTL IYYDYTDV 51 AGGACTAGAA GGTGGTTGGC AACTGATCGG GACTCAGACA ATGAAGCAAA CTCTGTTAAG CTGGAGGAAA TTGGGAGGAAA TTGAAGTCCG TATTAAAAAA AAGGAGTCCT TATTAAAAAA AAGGAGTCCT CTAACTATTA ATTTTGGTTTT GGCCTCAAGC CTGCACCGG AGTTGTTACA AGCAGTCAC GTGATTGTTACA GCCTCAAGC CTGCACCGG AGTTGTTACA TTGAGTTACA TTGAGTTACA TTGAGAGATAC TTAAAATTGTA TTGTTTTCCCA TTAAGCTTTA TTTGTTTTTT	180 240 300 60 120 180 240 360 420 780 960 960 1020 1080 1140 1260 1320 1380 1440 1560 1560 1620 1680 1740 1800	
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50 55 60 65 70 75	FYANMYTSIV NGQPTEDNIH ISQSSRKRKH PLSACNVCLD Seq ID NO: Nucleic Ac: Coding sequence GTTCGGCGCC CGAAAGGAGT AAGGCCGCGG GCAAGGAGT AAGTCTGGAG GTTCTGGAG GTTCTGGAG GTTCTGGAG GTTCTGGAG GTCTAGATAT GAGACTTGT TGAAGGTTT TGAAGGTTT TGAAGATTCT CTCTCTGTA AAATAGCCAG AGGCACATC CTCTCTGTA AGACTTCT TGGTTTTTTTTGTGCAG TATTTGGGAA CTTTGGGAA CTTTGGGAA CTTTGTGGCA TATTTGGGAA CTTCTGGGAA TCCCAGTTATT TCCCAGTTATT TCCCAGTTTAT TCCCAGTTTAT TCCCAGTTTAT TCCCAGTTTAT TCCCAGTTTAT TCCCAGTTTAT TCCCAGTTTAT TCCCAGTTTAT TCCCAGTTTAT TCCCAGTTTAT TCCCAGTTTAT TCCCAGTTTAT TCCCAGTTTAT TCCCAGTTTAT TCCCAGTTTAT TCCCAGTTTAT TCCCAGTTTAT TCCCAGTTTAT TCCCAGTTTAT TCCCAGTTAT CCAGTT TCCCAGT TCCCA	FLGLISIDRY DCSKLKSPLG NQSIRVVVAV PIIYFFMCRS 153 DNA see id Accession lence: 149- 11 AAAGCGCGGA GAGGGCCGA GAGTGGGAAG GCGCGCCG GAGATGAAAG GCGTATAATTATA GCATATACAC GACTATTACAC GACTATTACAC CTTCATGAG CTCACTCT TTAAGATAAC TTTTTAATG GACTGTCC TTTTTAATG GACTGTCC TTTTTCTAAT GACTGTCC TTTTTCTAAT GACTGTCC TTTTTCTAAT GACTGTCTC TTAAATTATAAT CCTTGTTCC TTTTTCTAAT GCAAGAGAAG AGCTGAATT TAAATGAAAG AGCAGAGAAG GCTGATTTTTCAA TGGGTGATC GACTTTTTAAT TAATTTTTCAG GCAAGTTTTTTAAT TCGACCGT GGAGGCTCTTT TATGACCCGT GGAGTCTTGT TCTATCCCCT TCTATCCCCT TCTCCCCT TTTTCCCCT TCTCCCCT TCTCCCCCT TCTCCCCT TCTCCCCCT	LKVVKPFGDS VKWHTAVTYV FFTCFLPYHL FFTCFLPYHL FFRRLFKKSN Quence 1 #: D80008 739 21 21 3 GCGGAGGCCG GAGCCCAGAT CGTCGGCAT AAGGGCACAC CTTTGTATGA TGATACAAC CATACCTGTA TGCAAATGC AAAGATCTCT TAGATAGAA AATTTGAAGT CTCGATGGAA AATTTGAAGT CTCAATGGAA AATTTGAAGT CTCAATGGAT AAGCTGTAT CTCAAGGTGTT AAGCTGTAT AAGCTGTAT CTAAACTGT TTAGACTG CTAAAGTGTT AAGCTGTAT AAGCTGTAT CTGAAGATACA TAAATATAAT ACATGGATTT ACCAGTATCC GGTATGTTT TTTTTTTTTT	RMYSITFTKV NSCLFVAVLV CRIPFTFSHL IRTRSESIRS 31 31 31 AGGCGAGAGG ACCATTTGG GTTCTGCGAA GCCTGCCTTC ACAAAACAG TATCAAATT TGACCGCTTG ATTACGATTT TGACCGCTTG ATTACGATTT TGACCGCTTG ATTACGATTT TGCTACTACT ACCACCAAAA TGATGATGGC CTTCACCTCC TGGCTAGGAA TCTTCACTAC CTTCACTAC CTAGCTAGT GAGATCACA TGATTTGTA GTTTAATGAC GTGTATTTCAAA GTGTTTTTCAAA GTGTTTTTCAAA GTGTTTTTCAAA GTGTTTTTCAAA GTTTAATGAC TCACATTTC CCACTTTGGA AAACAGCTGA AAACAGCTGA AAACAGCTGA TGGGTGTTGC CTTCTAGAAG GTTTTTTTTAAAG GTTTTTTTTAAAG GTTTTTTTT	LSYCWYIMA ILIGCYIAIS DRILDESAQK LQSYRRSEVR A1 CTGGCGCTGT CGTGAGAGCT CGTGAGAGCT TCTGGGTCA AAGCCATGGT CTGGGTCA CACATGGTA ACTCCT CTGATCGTA ACTTCAGTCC CTGATCAGTC CTGATCAGTC CTGATCAGTC CTGATCAGTC CTGATCAGTC CTGTATCAGTC CTGTATCAGTC CTCTTTGATT GTATAAATTT GTCTTCTGGTCAT TCCTGGTCAC CTCTTTAAT TCCTGGTCAT TCCTGTGAT ACATGCAA ATACTAATTT CACCATGGTG TCGAGAGAA ATACTAATTT CACCATGGTG TCGAGAAA CATTTAAAT ATCCGAGAAA ATTTTAAAT TCTGAGCAT GCAGTGGCGT TCTCTAGGCTC CCTCTTTAAAT TTCACATGCAC TTGACTGAAA ATACTAAATT TCACGAGAAA CATTTTAAAT TCTCAGCCTC TCTCTAGCCTC TCTCAGCCTC	VISLENTILIT RYIHKSSROF ILYYCKETTL IYYDYTDV 51 AGGACTAGAA GGTGGTTGGC AACTGATCGG GACTCAGACA ATGAAGCAAA CTCTGTTAAG CTGGAGGAAA TTGGGAGGAAA TTGAAGTCCG TATTAAAAAA AAGGAGTCCT TATTAAAAAA AAGGAGTCCT CTAACTATTA ATTTTGGTTTT GGCCTCAAGC CTGCACCGG AGTTGTTACA AGCAGTCAC GTGATTGTTACA GCCTCAAGC CTGCACCGG AGTTGTTACA TTGAGTTACA TTGAGTTACA TTGAGAGATAC TTAAAATTGTA TTGTTTTCCCA TTAAGCTTTA TTTGTTTTTT	180 240 300 120 180 240 360 420 480 540 660 660 720 780 840 900 960 1020 1140 1200 1320 1380 1440 1500 1560 1620 1680 1740 1860 1740 1860 1740 1860 1960 1960 1960 1960 1960 1960 1960 19

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WO 02/086443

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Seq ID NO: 181 DNA sequence Nucleic Acid Accession #: Eos sequence

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PCT/US02/12476 WO 02/086443

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                                                                                                          960
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                                                                                                        1080
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                                                                                                        1140
75
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         Nucleic Acid Accession #: EOS sequence
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5	CGGCGAGGGG	CCGCAGACCG	CCCCCATTCA	CCTAATCCAT	ACTACAGACA	ACAGAGAAAA	240
5		3/77 PYPY COUNTY OF CO.	CTCCTATACA	GGAGCACTGA	ATCAMMAN.	TOGGGWWWGW	300
		A TOTAL A TOTAL CO.	<u> </u>	CTCCTATCAA	TATTGALGAA	GAICIIACAC	360
	mman	CARTCTTARG	TALATTOLA	TTCAGGGTTG	GGATAAAACA	TCATIGGAAA	420
		TO STATE OF THE ST	CCCAAAACAG	TOGDDATTAA	TCTCACTAAL	GACTACCGIG	480
10		* COMMONICS CL 2 2	አጥርያርጥርያጥጥላ	AAGCAAGCAA	GATAACITII	CWCIGGGGVV	540
			CONTONCIO	ATAGTTTAGA	ALGACAAAAA	TITCCMCTIG	600 660
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	O > COO O CO O O	CTCTCN NACC	CACTTGGCTG	TTTTTTGTGA	AGTTCTTACA	ATGCAACAAT	960
		CATCCTCATC	CACTACTTAC	AAAACAATTT	TCGAGAGCAA	CAGTACAAGI	1020
			サンカイ カイカイ ひんかん	GAAAGGAAGA	GATTCATGAA	GCAGTITGIA	1080
20		TOTAL A A A COMP	CAGGCTGACC	CAGAGAATTA	TACCAGCCTT	CITGITALAI	1140
		THE PROPERTY OF THE PARTY OF TH	ጥስጥርስጥልሮሮል	TGATTGAGAA	GTTTGCAGTT	TIGIACCAGC	1200
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	GTGCTATTCT	CAATAATTTG	CTACCCAATA	TGAGTTATGT	TCTTCAGATA	CCTACCCATAL	1380
0.5	GCACTAATGG	CTTATATGGA	AAATACAGCG CCTGAATTAA	ACCAACTGAT	TGTCGACATG	AAGGAGGAGG	1440
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		MORROCCA 3/7	スペヤススペペペスカヤ	CCCCAACAAG	AGGAAGTGAA	TTCTCTGGMM	1620
		MOOGEN A MACA	ידידי על עיניידיינייטיי	CCACTTCCCA	ACCAGICACI	AAAIIAGCCA	1680
30			* CTTTTTTT	CTGTGACTGA	ACTUCCACCI	CHCMCIGIGG	1740
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70		* *************	™ ማእጥርምል ሮ ሞቸ	2 AAGTGAGGC	A CTTTCAGTG	I CCTAAATGGC	4140
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75			~ **************	~ ттстсасса	C AAGGCAGGA	A GAGAATCCAT T GAGAGCTTAG	4440
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         Protein Accession #: EOS sequence
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85	ACCOMON	AAAAAAAAA	GCATGAGTTC	D D D TO CO TO TO TO	CLTALCACACA	ССДУСССУУД	9000
0.7	AAAAA	AAAAACAAGG	GCWIGWGIIC	WWIGGETTH	~~~~~~~	COLOGICANI	2000
				3 CC3 CC3 CC-		THE COMMENT	0000
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WO 02/086443
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         Coding sequence: 1-394
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50	1   GAGCAACCTC CGACCCAGAG GCGGGGCCCA ACCTGCCACC GCTGTTGGGC GCCCCAGTGG CGAGGGGGTT TGACTCCTTG CATCCTCTG CTTGGAAGAC TCTTGCAGGT ATTCTATGAC	11   AGCTTCTAGT CTTCTCCAGC GCCACCTTCC CCTGAGCCAG TTCATTCTCG AGGATTTACT TGGATGTCCT CTGAATCTGA GATGAGGTGC CTGGCTATTT CCTATGACCCC	21   ATCCAGACTC GGCGGCGCAG GGAGTCCGGG CCTTCCTGGG CCTTCCTGGG CCTTCCTGGG GCAGCACATT CAATCTTTGT AGAAGATGAG TAGTTGCCAC CAGTCAATGC	CAGCGCCGCC CCAGCAGGGC TTGCCCACCT CCAGCGAGTC CGACAACATC GAGCACCGGG GCAACCACCGTT GATGGCTGTC AGCATGGTAT CAGGTACGAA	CCGGGCGCG TCCCCGCCTT GCAAACTCTC ATGGCCAACG GCCATCGTCA GTGACCGCCC CAGATCCAGT CGTGCCTTGA GGCATGAAGT ATTGGGGTG GGCAATAGAA TTTGGTCAGG	ACCCCAACCC AACTTCCTCC CGCCTTCTGC CGCGCTGCA GCACTGCCT AGGCCATGTA GCAAAGTCTT TGGTGGTTGG GTATGAAGTG CGATATTTCT TCGTTCAAGA CTCTCTTCAC	120 180 240 300 360 420 480 540 600 660 720
50 55	1 	11   AGCTTCTAGT CTTCTCAGC CCTGAGCCAG TTCATTCTCG AGGATTTACT TGGATGTCCT CTGAATCTGA GGATGATAG GATGAGGTGC CTGCTATTACCC CCTGCTATTACCC GCTGCTTCTC	21   ATCCAGACTC GGCGGCGCG GGAGTCCGG CCTCCTGGG CCTATGCCGG CCTATGCCGG GCAGCACATT CAATCTTTGT AGAAGATGAG TAGTTGCAC CAGTCAATGC CAGTCAATGC TCTGCCTTCT	CAGCGCCGCC CGAGCAGGGC TTGCCACCT CGAGCGAGTC ATGGATCGGC CGACAACATC GAGCACCGGG GCAACCATC GATGGCTGTC AGCATGGTAT AGCATGGTAT CAGGTACGAA GGGAGGTACCA	CCGGGCGCG CCAAACTCTC ATGGCAACG GCCATCGTCA GTGACCGCCC CAGATCCAGT CGTGCCTTGA GCCATGAGT ATTGGGGGTG GCAATAGAA TTTGGTCAGG CTACTTTGCT	ACCCCAACCC AACTTCCTCC CGCCTTCTGC CGCGGCTGCA GCACTGCCCT AGGCCATGTA GCAAAGTCTT TGGTGGTTGG GTATGAAGT CGATATTTCT TCGTTCAAGA CTCTCTTCAC GTTCCTGTCC	120 180 240 300 360 420 480 540 600 720 780
50	1   GAGCAACCTC CGACCCAGAG GCGGGGCCCC GCTGTTGGGC GCCCCAGTGG GCGAGGGGCTG TGACTCCTTG CATCCTCCTG CTTGGAAGAG TCTTGCAGGT ATTCTATGAC TGGCTGGGCT CCGAAAAACA GAAAGACTAC	11	21   ATCCAGACTC GGCGGCGCAG GGAGTCCGGG CCTTCCTGGG CCTATGCCGG GCAGTCAGCAGAT CAATCTTTGT AGAAGATGAG TAGTTGCCAC CAGTCAATGC TCTGCCTTCCTCAACACCAAG GAGGCAAAAAG GAGGCAAAAAAG	CAGCGCCGCC CCAGCAGGGC TTGCCCACCT CGAGCAGGTC CGACAACATC GAGCACCGGG GCCACCGTT GATGGCTGTC AGCATGGTAT CAGGTACGAA GGGAGGTGCC GCCCTATCCA GGAGAAAATCA	CCGGGCGCGG CCCGCCTT GCAAACTCTC ATGGCCAACG GCCATCGTCA GTGACCGCC CAGATCCAGT CGTGCCTTGA GGCATGAAGT ATTGGGGGT GGCAATAGAA TTTGGTCAG CTACTTTGCT TAAACCTGCAC TGTTGAAACA	ACCCCAACCC AACTTCCTCC CGCCTTCTGC CGCGCTTCTGC CGGGGCTGCA GCACAGTTA GCAAAGTCTT TGGTGGTTGG GTATGAAGTG CGATATTTCT TCGTTCAAGA CTCTCTCAC GTTCCTGTCC CTTCCAGCGG AACCGAAAAT	120 180 240 300 360 420 480 540 600 660 720
50 55	1   GAGCAACCTC CGACCCAGGG GCGGGGCCA ACCTGCCACC GCTGTTGGGC GCCCAGTGG GCACCAGTGG CATCCTCCTG CATCCTCCTG CATCCTCTG CATCTCAGGT ATTCTATGAC TGGCTGGGCT CCGAAAACA GAAAGACTAC GGACATTGAG	11   AGCTTCTAGT CTTCTCAGC CCTGAGCCAG CCTGAGCCAG TTCATTCTCG AGGATTACT TGGATGTCCT CTGAATCTGA GGATGATAG GATGAGGTGC CTGGCTATTT CCTATGACCC GCTGCTTCTC ACCTCTTACC GTGTGACACA ATACTATCAT	21   ATCCAGACTC GGCGGCGGG GGAGTCCGGG CCTATGCCGG CCTATGCCGG CCATGCCGA CAGCACATT CAATCTTTGT AGAAGATGAG TAGTGCAC CAGTCAATGC CAGTCAATGC CTGCCTTCT CAACACCAAG GAGGCAAAAG TAACATTAGG	CAGCGCCGCC CCAGCGAGGC TTGCCACCT CCAGCGAGTC ATGGATCGGC CGACAACATC GAGCACCGGG GCACACCGTT GATGGCTGTC AGCATCGATCACCACGGTAGCAACCACCGTT GATGGCTGTC GAGGAACATC GGGAGGTACCA GGGAAGAATCA ACCTTAGAAT	CCGGGCGCGG CCAACCTCT GCAAACTCTC ATGGCCAACG GCCATCGTCA GTGACCGCCC CAGATCCAGT ATTGGGGTG ATTGGGGGTG GGCAATAGAA TTTGGTCAGG CTACTTTGCT AAACCTGCAC TGTTGAAACA TTTTGGAACT TTTTGGAACT TTTTGGAACT TTTTGAAACA TTTTGGAATT	ACCCCAACCC AACTTCCTCC CGCCTTCTGC CGCGCTTCTGC CGCACTGCCCT AGGCCATGTA GCAAAGTCTT TGGTGGTTGG GTATGAAGTG CGATATTTCT TCGTTCAAGA CTCTCTTCAC GTTCCTGTCC CTTCCAGCGG AACCGAAAAT GTAATCTGAA	120 180 240 300 360 420 480 540 660 720 780 840 900 960
50 55	1   GAGCAACCTC CGACCCAGGG ACCTGCACC GCTGTTGGGC GCCCCAGTGG GCCCCAGTGG CGAGGGGCTG TGACTCCTTG CATCCTCTG CATCTCTGAGGC TCTTGCAGGT ATTCTATGAGC TGGGTGGGCT CCGAAAAACA GAAAGACTAC GGACATTGAG GTATGGTATT	11   AGCTTCTAGT CTTCTCCAGC CCCAGCCAG CCCACCTTCG CCTGAGCCAG TTCATTCTCG AGGATTTACT TGGATGTCCT CTGAATCTGA GATGAGGTGCT CTGATCTCC CTGCTATTT CCTATGACC GCTGCTTCTC ACCTCTTACC GTGTGACACA ATACTATCAT ACAAAACAAA	21   ATCCAGACTC GGCGGCGCAG GGAGTCCGGG CCTATCCTGGG CCTATCCTGGG GCAGCACATT CAATCTTTG AGAAGATGAG TAGTTAGCAC CAGTCAATGC CAGTCAATGC CAGTCAATGC CAGTCAATGC CAGTCAATGC CAGTCAATGC CAGTCAATGC CAGTCAATGC CAGTCAATGC CAACACAAC CAGCAAAAG CAACAAACAA	CAGCGCCGCC CGAGCAGGCC TTGCCACCT CGAGCGAGTC ATGGATCGGC CGACAACATC GAGCACCGGG GCAACCATC GAGCACCGTT GATGGTATCGA GGGAGTACGAC GGGAGCAGCGT CAGGTACGAA GGGAGGTACCA GAGAAAATCA AACCCAT AAAAACCCAT	CCGGGCGCGG CCAAACTCTC ATGGCAACG GCAATCGTCA GTGACCGCCC CAGATCCAGT CGTGCCTTGA ATTGGGGTG GCAATAGAA TTTGGTCAGG CTACTTTGCT AAACCTGCAC TTTGGACAC TTTTGGACAC TTTTGGTAAACA TTTGGGTATT GTGTTAAAACT GTGTTAAAAT	ACCCCAACCC AACTTCCTCC CGCCTTCTGC CGCGGCTGCA GCACTGCCCT AGGCCATGTA GCAAAGTCTT TGGTGGTTGG GTATGAAGTG CGATATTTCT TCGTTCAAGA CTCTCTTCAC GTTCCTGTCC CTTCCAGCGG AACCGAAAAT ACTCAGTGCT	120 180 240 300 420 480 540 600 720 780 840 900 960 1020
50 55 60	1   GAGCAACCTC CGACCCAGAG GCGGGGCCCA ACCTGCCACC GCTGTTGGGC GCCCCAGTGG GCAGGGGCTG TGACTCCTTG CATCCTCTG CTTGGAAGAC TCTTGCAGGT ATTCTATGAC TGGCTGGGCT CCGAAAAACA GAAAGACTAC GGACATTGAG GTATGGTATT AAACATGGCT	11   AGCTTCTAGT CTTCTCCAGC CCTGAGCCAG TTCATTCTCG AGGATTTACT TGGATGTCTC CTGAATCTGA GAGGGATAGG GATGAGTGC CTGGCTATTT CCTATGACCC GCTGCTTCTC ACCTCTTACC GTGTGACACA ATAAAACAAA TAATCTTATT	21   ATCCAGACTC GGCGGCGCAG GGAGTCCGGG GCAGGCCC CCTTCCTGGG CCTATGCCGG GCAGCACATT CAATCTTTGT AGAAGATGAG TAGTTGCCAC CAGTCAATGC CTCCTCCTTCC TCACACACAAG GAGGCAAAAG TAACATTAGC TAACACTAAG TAACATTAGC TTATCTTCTT	CAGCGCCGCC CGAGCAGGGC TTGCCCACCT CGAGCGAGTC GCGACACATC GAGCACCGG GCAAGCAACCT GATGGCTGT GATGGCTGT AGCATGGTAT CAGGTACGAC GCCTATCCA GCCTTATCCA ACCTTAGAAT TCCTCAATAT	CCGGGCGCGG CCAAACTCTC ATGGCCAACG GCCATCGTCA GTGACCGCC CAGATCCAGT CGTGCCTTGA GGCATGAAGT ATTGGGGTG GGCATAGAA TTTGGTCAGA CTACTTTGCT AAACCTGCAC TGTGAAACA TTTGGGTATT GTGTTAAACA AGGAGGGAAA	ACCCCAACCC AACTTCCTCC CGCCTTCTGC CGCGGCTGCA GCACTGCCCT AGGCCATGTA GCAAAGTCTT TGGTGGTTGG GTATGAAGT CGATATATTT TCGTTCAAGA CTCTCTTCAC GTTCCTGTCC CTTCCAGCGG AACCGAAAAT GTAATCTGAA ACTCAGTGCT ATTTTACCAT	120 180 240 300 420 480 540 600 720 780 840 900 960 1020
50 55	GAGCAACCTC CGACCCAGG GCGGGGCCCA ACCTGCCACC GCTGTTGGGC CGCCCAGTGG CGACCCGTGC CATCCTCCTG CATCCTCTG CATCCTCTG ATTCTATGAC TCTTGCAGGT ATTCTATGAC GAAAACAA GAAAGACTAC GGACATTGAG GTATGGTATT AAACATGGCT TTGTATTACT	11   AGCTTCTAGT CTTCTCAGC GCCACCTTCG CCTGAGCCAG TTCATTCTCG AGGATGTCCT TGGATGTCCT CTGAATCTGA GGATGATAG GATGAGGTGC CTGGCTATTT CCTATGACCC GCTGCTTCTC ACCTCTTACC GTGTGACACA ATACTATCAT ACAAAACAAA	21   ATCCAGACTC GGCGGCGGG GGAGTCCGGG CCTATGCCGG GCGTGTCGCA GCAGCACATT CAATCTTTGT AGAAGATGAC TAGTGCAC CAGTCACTC CAGCACTAT TAGACACAAG GAGGCAAAAG TAACATTAGG CAAACAAACA TTATCTTCT GAGTAATCCT TAGTTCTT GAGTAATCAT	CAGCGCCGCC CGAGCAGGC TTGCCCACCT CGAGCAGTC CGAGCAACATC GAGCACCGGG GCAACAACATC GAGCACCGTT GATGGCTGTC AGCATGGTAC AGCATGGTAC AGGAGTACC AGGAAGAAC GGGAGGTGC GCCCTATCCA AGCAAACAT AAAAACCCAT TCCTCAAATG	CCGGGCGCGG CCAAACTCTC ATGGCAACG GCCATCGTCA GTGACCGCCC CAGATCCATGA GGCATGAAGT ATTGGGGTTGA GGCAATAGAA TTTGGTCAGG CTACTTTGCT AAACCTGCAC TGTTGAAACA TTTGGGTATT GTGTTAAAAT GGGAAGGGAA	ACCCCAACCC AACTTCTCC CGCCTTCTGC CGCGCTTCTGC CGCGCTTCTGC CGCCATGTA GCACAGCCT TCGTGGTTTG GTATGAAGTG CGATATTTCT TCGTTCAAG CTTCCTTCAC CTTCCAGCGG AACCGAAAAT GTAATCTGAA ACTCAGTGT ATTTACCAT GCTCCTTAAA	120 180 240 300 420 480 540 600 720 780 840 900 960 1020
50 55 60	1   GAGCAACCTC CGACCAGAG GCGGGGCCCA ACCTGCCACC GCTGTTGGGC GCCCAGTGG GCAGGGGCTG TGACTCCTTG CATCCTCTG CATCCTCTG CATCCTCTG CATCCTGAGAC TCTTGCAGGT ATTCTATGAC TGGCTGGGCT ACCGAAAACA GAAAGACTAC GACATTGAG GACATTGAG TATGGTATT TATATATAGA CTCATTATGT	11   AGCTTCTAGT CTTCTCCAGC CCGAGCCAG CCTGAGCCAG TTCATTCTCG AGGATTTACT CTGAATCTGA GAAGGATAGG GATGAGTACC CTGGCTATTT CCTATGACCC GCTGCTTCTC ACCTCTTACC GTGTGACACA ATAAACAAA TAATCTTATT GCTTCCCATT TATGTATATA TGGTACTAGC	21   ATCCAGACTC GGCGGCGCAG GGAGTCCGGG CCTTCCTGGG CCTATGCCGG GCAGCACATT CAATCTTTGT TAGAAGATGAG TAGTTGCCAC CAGTCAATGC TCACCACAG GAGGCAAAAG TAACATTAGC TAACACAAAG GAGGCAAAAG TAACATTTGT GAGTAATCTT GAGTAATCATT ATACTTTAAAA	CAGCGCCGCC CGAGCAGGGC TTGCCCACCT CGAGCAGGGC CGACAACATC GAGCACCGGG GCCACCGTT GATGGCTGTC GAGTACGAA GGGAGGGGT CCCTATCCA ACCTTAGAAT AAAACCCAT TCCTCAAATAT ACTCAAATGA TCTTATAAAA TATCTCTAAA	CCGGGCGCGG CCAACCTT GCAAACTCTC ATGGCCAACG GCCATCGTCA GTGACCACG CCAGATCCAGT CGTGCCTTGA GGCATGAGT ATTGGGGGTG GGCATAGAA TTTGGTCAGA CTACTTTGCT TAAACCT TGTTGAACCA TTTGGGTATT GTGTTAAACT AGGAGGGAAG GGGAAGGGGT ATAGGTAAACT ATAGACAGTA ATAGGTAAACT	ACCCCAACCC ACCTCTCTC CGCCTTCTGC CGCGCTTCTGC CGCGCTTCTGC CGCGCTGCA AGGCCATGTA GCAAAGTCTT TGGTGGTTGG GTATGAAGTG TCGTTCAAGA CTCTCTCAC GTTCCTGTCC CTTCCAGCGG AACCGAAAAT GTAATCTGAT GCTCTTAAC ATTTACCAT GCTCCTTAAC AAATACTATT GTATTTAATT	120 180 240 360 420 660 720 780 840 900 1020 1080 11400 1260
50 55 60	1   GAGCAACCTC CGACCCAGAG GCGGGGCCCA ACCTGCCACC GCTGTTGGGC GCCCCAGTGG GCACGGGCT TGACTCCTTG CATCCTCTG CATCCTCTG CTTGGAAGAC TCTTGCAGGT ATTCTATGAC TGGAAAACA GAAGACTAC GGACATTGAG GTATTGGTATT AAACATGGCT TTGTATTACT TATATATACT TATATATAGA CTCATTATGT	11   AGCTTCTAGT CTTCTCCAGC GCCACCTTCG GCCACCTTCG CCTGAGCCAG TTCATTCTCG AGGATTTACT CTGAATCTCG GAGTGATACG GATGACTCG CTGGCTATTT CCTATGACCC GCTGCTTCTC ACCTCTTACC GTGTGACACA ATACTATCAT ACAAAACAAA	21   ATCCAGACTC GGCGGCGCAG GGAGTCCGGG CCTTCCTGGG CCTATGCCGG GCAGCACATT CAATCTTTGT AGAAGATGAG TAGTGGCAC CAGTCAATGC TAGCCAC CAGTCAATGC TAACACCAAG CAACCAAG CAACCAAG CAACCAAG TAACATTAGG TAACATTAGG TAACATTATT TAACATTATAA ATTGGTATAT	CAGCGCCGC CCAGCGAGGC TTGCCCACCT CGAGCGAGTC CGACAACATC GAGCACCGGG GCACACCGTT GATGGCTGTA ACCACCGTT AGGATCGGA GGGAGGTGCC GCCCTATCCA ACCTTAGAAT ACCTTAGAAT ACAAACCCAT ACTCAAATG TCCTCAAATG TCTTATAAAA TATCTCTATTTC	CCGGGCGCGG CCAAACTCTC ATGGCAACG GCCATCGTCA GTGACCGCCC CAGATCCAGT ATTGGGGTG ATTGGGGTG ATTGGGGTG ATTGGGGTG ATTGGTCAGG CTACTTTGCT AAACCTGCAC TTTTGGAACA TTTGGGTATT GTGTTAAAAT ATGGGAAG GCGAAGGGGT ATAGACAGTA ATAGACAGTA ATAGACAGTA ATAGACAGTA ATAGACAGTA ATAGACAGTA ATAGACATAT GTCCTTATAT	ACCCCAACCC AACTTCTCC CGCCTTCTGC CGCGCTTCTGC CGGGGCTGCA AGGCCATGTA GCAAAGTCTT TGGTGGTTGG GTATGAAGTG CTTCTCAC GTTCCAGCG CTTCCAGCG CTTCCAGCGAAAAT GTAATCTGAA ACTCAGTGCT ACTCAGTGCA ACTCAGTGCT GCTCCTTAAA AAAATACTAT GTATTTAACTA	120 180 240 300 360 420 540 600 720 780 960 1020 1080 1140 1260 1320
<ul><li>50</li><li>55</li><li>60</li><li>65</li></ul>	GAGCAACCTC CGACCCAGGG GCGGGGCCCA ACCTGCCACC GCTGTTGGGC GCCCCAGTGG CGAGGGGCTG CATCCTCCTG CATCCTCTG CATCCTCTG CATCTATGAGAC TCTTGCAGGT ATTCTATGAC GGACATTGAG GGACATTGAG GTATGGTATT AAACATGGCT TTGTATTACT TTGTATTACT TATATATAGA CTCATTATGT CCATTATGT CCATTATGT CAGTCAAATA	11   AGCTTCTAGT CTTCTCCAGC CCTGAGCCAG CCTGAGCCAG TCCATTCTC TGGATGTCCT TGGATGTCCT CTGAATCTGA GGAGTGATAC CCTGCTATTC CCTATGACCC CCTGCTATTT CCTATGACCC ACCTCTTACC GTGTGACACA ATACTATACAT ACAAAACAAA	21   ATCCAGACTC GGCGGCGCGC GGAGTCCGG GCGGGCGCC CCTTCCTGGG CCTATGCCGG GCAGTACACT CAATCTTTGT AGAAGATGAG TAGTCACTC CAACCAAG GAGGCAAAAG GAGGCAAAAG TAACATTAGT CAACACAAG GAGGAAAACA TTATCTTCTT AGAGAATCAT TACATGTTTT ATACTTAAAA ATTGGTATTA TTCTTCATTA	CAGCGCCGC CCAGCAGGGC TTGCCACCT CCAGCAGGGC ATGGATCGGC CGACAACATC GAGCACCGGG GCACACGTT GATGGCTGTC AGCATCGAC AGCAACAT AGGATCGGC GCCCTATCCA GGGAAAATA AACCCAT TCCTCAATAT ACTCAAATGG TCTATTAAAA TATCTCTAAAA TATCTCTAAAA TTTCTTTTTC GCTTTGGGTG	CCGGGCGCGG CCAACCTCT ATGCCAACG GCCATCGTCA GTGACCGCCC CAGATCCAGT CGTGCCTTGA ATTGGGGGTG GCCAATAGAA ATTTGGTCAGG CTACTTTGCT AAACCTGCAC TGTTGAAAC TTTTGGAAC TTTTGGAAC TTTGGGAAT AGGAACA TATGAACA TATGAACA TATGAACA TATGAACA TATAAAT AGGAGGGAA ATAGGTAAT ATAGGTAAT CCTTTGCCAC	ACCCCAACCC AACTTCCTCC CGCCTTCTGC CGGGGCTGCA GCACTGCCCT AGGCCATGTA GCAAAGTCTT TGGTGGTTGG GTATGAAGA CTCTCTTCAC GTTCCTTCAC GTTCCAGCGG AACCGAAAAT ACTCAGTGCT ATTTTACCAT ACTTCACAAAA AATACTATT GTATTTAATT ACATATGTAA AAGACCTAGC	120 180 240 300 360 420 540 600 720 780 840 900 960 1020 1140 1260 1260 1320
50 55 60	1   GAGCAACCTC CGAGCCAGC GCGGGGGCCCA ACCTGCCACC GCTGTTGGGC GCACCCAGTG TGACTCCTTG CATCCTCTG CATCCTCTG CATCCAGAGAC CTGAGAGAC TGACTGAGGT ATTCTATGAC GGACATTGAG GTATGGATTATATACATACATTACAT	11   AGCTTCTAGT CTTCTCCAGC CCGAGCCAG CCTGAGCCAG TTCATTCTCG AGGATTTACT TGGATGTCCT CTGAATCTGA GAGTGATAG GATGAGTATT CCTATGACCC GCTGCTTCTC ACCTCTTACC GCTGCTTCTC ACCTCTTACC GTTGACACA ATACTATCAT TCATACATATT TGGTACATATT TGGTACATATT TGGTACATATT TGGTACATATT TGGTACATATT TGGTACATATT TGGTACATATT TGATACTATCAT TCATTTACTC AAGGATGATT TCATTTACTC AAGGATGATAT	21   ATCCAGACTC GGCGGCGCAG GGAGTCCGGG GCAGGCGCC CCTTCCTGGG GCAGCACATT CAATCTTTGT AGAAGATGAG TAGTGCAC CAGTCAATGC CAGTCAATGC CAGTCAATGC CAGTCAATGC CAACAACA TAACATTAGG CAAACAAACA TTACTTCTT GAGTAATCAT TACATGTTTT ATACTTAAAA ATTGGTATAT TTCTTCAATT	CAGCGCCGCC CAGCGAGGGC TTGCCCACCT CGAGCAGGGC ATGGATCGGC CGACAACATC GAGCACCGT GATGCATCGT GATGCTGTC GAGCACGGT GATGGTATCAA GCATGGTATCAA ACCTTAGAAT AAAACCCAT TCCTCAATAT ACTCAATAT ATTCTTTTCC CTTTGGGTG CTTTCGTGTG CTTTTGGGTG CTTCATGGGTG CTTCATGGGTG CTTCATGGGTG CTTCATTGGTG CTTCATTGGTG CTTCATTGGTG CTTCATTGGTG CTTCATTGGTG CTTCATGGGTG CTTCATGGGTG CTTCATGGGTG CTTCATGGGTG CTTCATGGGTG CTTCATGGGTG CTTCATGCGTT	CCGGGCGCGG CCAAACTCTC ATGGCCAACG GCAATCGTCA GTGACCGCC CAGATCCAGT CGTGCCTTGA GGCATCAGT GTGCTTGA GGCATGAGT CTTGGTCAGG CTACTTTGCT AAACCTGCAC TTTGGTATAACA TTTGGTATT AGGAGGGAGGGT ATAGGAAGT ATAGGAAGGGAAGGGT ATAGGAAGTA ATAGGTAAAT ACGTAAACT CCTTTATAT CCTTTTGCT CCCCTTTTCA	ACCCCAACCC AACTTCCTCC CGCCTTCTGC CGCGGCTGCA GCACTGCCCT TGGTGGTTGG GTATGAAGTCT TCGTTCAAGA CTCTCTTCAC GTTCCTGTCC CTTCCAGCGG AACCGAAAAT GTAATCTGAA ACTCAGTGCT ATTTTACCAT GCTCCTTAAC ACTCCTTAAC TGAATCTCAT ACTTTACCAT GCTCCTTAAC AAATACTAAT GCATATTTAACTAT ACATATTTAACTAT ACATATTTAAC TATATTTAACTAT ACATATTTAAC TATATTTAAC TATATTTAAC TATATTTAAC TATATTTAAC TATATTTAAC TATATTTAAC TATATTTAAC TATATTTAAC TATATTTAAT ACATATTTAAT ACATATTTAAT TATATTTAAT TATATTTAAT TATATTTATT	120 180 240 300 360 480 540 660 720 780 900 900 91020 1080 11200 1260 1320 1380 1440
<ul><li>50</li><li>55</li><li>60</li><li>65</li></ul>	GAGCAACCTC CGACCCAGGG GCGGGGCCCA ACCTGCCACC GCTGTTGGGC GCCCCAGTGG CGAGGGGCTG CATCCTCCTG CATCCTCCTG CATCCTCTG CATCCTCTG ATTCTATGAG TCTGCAGGT ATTCTATGAG GAAAACAA GGACATTGAG GTATGGTATT AAACATGGCT TTGATATTACT TATATATAGA CTCATTATGT CCATATTGT CCATATTGT CTATTTTTA	11   AGCTTCTAGT CTTCTCCAGC CCTGAGCCAG CCTGAGCCAG TCCATTCTC TGGATGTCCT TGGATGTCCT CTGAATCTGA GGAGTGATAG GCAGTGATAG CCTGCTATTC CCTGTATACC GCTGCTATTT CCTATGACCC ATCTTACC GTGTGACACA ATACTATACAT ACAAAACAAA	21   ATCCAGACTC GGCGGCGCG GGAGTCCGGG CCTATGCCGG CCTATGCCGG CCTATGCCGG CCTATGCTGCA GCAGCACATT CAATCTTTGT AGAAGATGAG TAGTCATTC CAACACAAG GAGGCAAAAC TTATCTTCT TAACATTATCT TACATTATAAA ATTGGTATAT TTCTTCATTA TCTTTCATTA TAGACTTA	CAGCGCCGC CCAGCGAGGGC TTGCCCACCT CCAGCGAGTC CGACAACATC GAGCACCGGG GCACACCGTT GATGGCTGTC AGCATCGGT GATGGCTGTC AGCATGGTAT CAGGTACCA GGAGAACATC GCCCTATCCA GAGAAACAT ACCTAGAAT ACCTAAATG TCTCAAATG TCTTAGAAT TTTCTTTTTC CCTTTGGGTG CTTCATGCTT CATGTATT ACAGTTATT ACATTTCATA	CCGGGCGCGG CCAACTCTC ATGCCAACG GCCATCGTCA GTGACCGCCC CAGATCCAGT ATTGGGGTG ATTGGGGTG ATTGGGGTG ATTGGTCA GGCAATAGAA ATTGGTCAGG CTACTTTGCT AAACCTGCAC TGTTGAAACA TTTGGTAAT GGGAAGGGGA ATAGGAAGA ATAGGTAAAT ACGAGGGAGA ATAGGTAAAT CCTTTGCCAC GCCCTTTTCA AAGCCCTTATA GCCTACATTT	ACCCCAACCC AACTTCCTCC CGCCTTCTGC CGCGCTTCTGC CGCGCTGCCCT AGGCCATGTA GCAAAGTCTT TGGTGGTTGG GTATGAAGTG CGATATTTCT TCGTTCAAC GTTCCTGCC CTTCCAGCGG AACCGAAAAT GTAATCTGAA ACTCAGTGCT ATTTTACCAT ACTATTTACTAA AAATACTATT GTATTTAATT TGATTTTATT ACATATGTAA AAGACTAGC TATGTTATA TGTATTTATT TGTATTTATT TTGTTTTTTTT	120 180 240 300 360 420 540 600 720 780 840 900 960 1020 1140 1260 1260 1320
<ul><li>50</li><li>55</li><li>60</li><li>65</li></ul>	GAGCAACCTC CGACCACCACC GCTGTTGGGC GCCCCAGTGG GCGGGGGCTG TGACTCCTTG CATCCTCTG CATCCTCTG CATCCTCTG ATTCTATGAC TCTTGCAGGT ATTCTATGAC TGGCTGGGCT GGACATGAGACATTACTATACAC GTATGGTATT AAACATGGCT TTGTATTACT TATATATAGA CTCATTATGT CCATATTGAT CAGTCAAATA CTAATTTACC TTATTTTTCATTGT TTCATTGGT TTCATTGGT TTCATTGGT TTCATTGGT TTTCATTGGT TTTCATTGGT TTCATTGGT TTCATTGGT TTCATTGGT TTCATTGGT TTCATTGGT TTCATTGGT TTCATTGGT TTCATTGGT	11   AGCTTCTAGT CTTCTCCAGC CCCAGCCTTCG CCCAGACCAG CCCAGACCAG CCCAGACCAG CCCAGACCAG CCCAGACCAG CCCAGACCAG CCCAGACCAG CCCAGACCAG CCAGACCAG CCAGACCAC CCAGACCAC CCAGACCAC CCAGACCAC CCAGACCAC CCAGACCAC CCAGACCAC CCAGACCAC CCAGACCAC CCAGACCAC CCAGACCAC CCACACCAC CCACACCAC CCACACCAC CCACACCAC	21   ATCCAGACTC GGCGGCGCAG GGAGTCCGGG CCTATGCCGG GCAGGCACATT CAATCTTTGT AGAGAATGA TAGTTGCAC CAGTCAATGC CAGTCAATGC CAGTCAATGC CAGTCAATGC CAGTCAATGC CAGTCAATGC TAGCATCT CAACACCAAG GAGGCAAAAG TAACATTAGG TAACATTAGG TAACATTATT TACATTAAAA ATTGGTATTT ATACTTCATTA TTCTTCAATT TCTTTCAATT TTGTAATT TGAGACTTAG TGAATCTAGA TAGACTTAG TGAATCTAGT TGAATCTACATT ATAGACTTAG TGAATCTAGAAA ATTGCATTA ATTGCATTA TTGAATCTAGAAA ATTAGAACTTAG TGAATCTAGAAACAAATCAGAACAAAATCAGAAC	CAGCGCCGCC CGAGCAGGCC TTGCCACCT CGAGCGAGTC ATGGATCGGC CGACAACATC GAGCACCGT GACACCGT GATGGCTGTC AGCATGGTAT AGCATGGAA AGGAAGCACC GCCACCGTT CAGGTACGAA AGGAAGCACC GCCATCCA AGAAAATCA AACCTATACAA TCCTCAAATAT TCTAAATAG TCTAAATAG TCTATATATA TATCTTTATC CGTTTGGGTG CATTGGTTATT ACATTATATA TATCTTTATT TTTGGAGCCA	CCGGGCGCGG CCAACCTCT ATGGCCAACG GCCATCGTCA ATGGCCACGC CAGATCCACG CCAGATCCACG CCAGATCCACG CCAGATCCACG CCAGATCCACG CAGATCCACG CAGATCCACG CAGATCCACG CAGATCCACG CAGATCCACG CAGATCCACG CAGATCCACG CAGATCCACG CTACTTTGCT AAACCTGCAC TTTGGACAA ATGGAGGAAG GGGAAGGGGT ATAGACAGTA ATAGGTAAAT CCTTACAAC CCCTTTTCA AAGCCCTTATC AAGCCCTTATC AAGCCCTTACT AATCTTTCTG	ACCCCAACCC ACCTTCTGC CGCGTTCTGC CGCGTTCTGC CGCGGTCTGCA CACATGCCCT AGGCCATGTA GCAAAGTCTT TGGTGGTTGG GTATGAAGT CTCTCTCAC GTTCCTGTCC CTTCCAGCGG AACCGAAAT ACTCAGTGCT ATTTTACCAT ACTCATCAAA ACTCAGTGCT ATTTTAACTAT GCTCCTTAAA AAATACTAAT GCTCCTTAAA AAATACTAAT TGATTTAATT ACAATATGTAA AAGACCTAGC TATTTTTTTTTT	120 180 240 300 360 480 540 660 720 780 900 900 1020 11200 1260 1320 1380 1440 1560 1560 1620
<ul><li>50</li><li>55</li><li>60</li><li>65</li><li>70</li></ul>	1   GAGCAACCTC GAGCCAGAG GCGGGGCCCA ACCTGCACC GCTGTTGGGC GCCCAGTGG GCAGGGGCTG TGACTCCTTG CATCCTCTG CATCCTCTG CATCCTCTG CATCAGAGAC ACTCATCAGAGAC ACAGAGACA GAAAGACAC GAAAGACAC GAAAGACATTGAG TATATATACT TATATATACT TATATATACT TATATATA	11   AGCTTCTAGT CTTCTCCAGC CCGAGCCAG CCTGAGCCAG TTCATTCTCG AGGATTTACT TGGATGTCTC CTGAATCTGA GAGTGATAG GATGAGTGC CTGGCTATTT CCTATGACCC GCTGCTCTTC ACCTCTTACC GTGTGACACA ATACTATCAT TATTATATAT TGATACTATAT TGATACTATAT TCATTTACT CAATTACT CAATTACT CAATTACT CAATTACT CAATTACT CAATTACT CAATTACT CCAATTACT CCAATTACT CAATTACT CAATTACT CAATTACT CCAATTACT CCCAATTACC CCTGTTGACC	21   ATCCAGACTC GGCGGCGCGC GGAGTCCGGG GGAGTCCGGG CCTTCCTGGG CCTTCCTGGG CCTATGCCGG GCAGCACATT CAATCTTTGT AGAAGATGAG TAGTTGCCAC CAGTCAATGC CAGTCAATGC CAGTCAATGC TCAACACAAG GAGGCAAAAG TAACATTACT TAACATTTTT ATACTTCTT GAGTAATCAT ATTCTCATTA ATTGGTATAT ATTCTTCAATT ATAGCACTG TGAATCAACA AATCAGAAC AAATCAGAAC AAATCAACA AATCAGAAC AAATCAACA AATCAGAAC AAATCAACA AATCAACA AATCAACA AATCAACA AATCAACA ATTCCACACA	CAGCGCCGC CGAGCAGGC TTGCCCACT CGAGCAGGC TTGCCCACCT CGAGCACATC GAGCACCGGG GCAACCATC GAGCACGGG GCAACCACGT GATGGCTGT ACAGTACACA GGGAGGTAC ACCTTAGAAT ACAAACAC ACCTTAGAAT ACAAATCA ACCTAATAT ACTCAAATGG TCTTATTAAAA TTTCTTTTTC GCTTGGGTG CTTCATGCGT CATCGTTATT ACATTCATA TTGGTGGTT ACATTTCATA TTTGGAGGC CATCGTTAC TTTGGAGGCA ATCCCTGTAC	CCGGGCGCGG CCAAACTCTC ATGGCCAACG GCCATCGTCA GTGACCGCCC CAGATCCATG ATTGGGCTA ATTGGGGTA ATTGGGGTA ATTGGGGTA ATTGGGTAA TTTGGTCAGG CTACTTTGCT AAACCTGCAC TTTTGAAACA TTTGGGTATT GTGTTAAACA TTTGGGTATT GTGTAAACA TTTGAAACA TTTGGTAAAC TTTGGTAAAC TTTGGTAAAC TTTTGAAACA TTTGGTAAAC GCGAAGAGAA GCGAAGAGAA GCGAAGAGAA GCCTTTTCA AACCCTTTTCA AACCCTTATT GCCTACATTT AATCTTTCTG TCTGACCCAT	ACCCCAACCC ACCTCTCTC CGCCTTCTGC CGCGTTCTGC CGCGTTCTGC CGCGTTCTGC CGCCTTCTGC CGCATGTTA GCAAAGTCTT TCGTTCATG CTATCTCACC GTTCCTTCAC GTTCCTTCAC GTTCCACCG AACCGAAAAT GTAATCTGAA ACTACTGTAC GTTCCTTAAA AAATACTATT GTATTTAACCAT GCTCCTTAAA AAATACTATT TCGTTTCATTTTTAATT ACATATGTTAA AAGACCTAGC TATGTTTTTTTTTT	120 180 240 300 360 420 540 600 720 780 900 900 91080 1140 1260 1320 1320 1340 1500 1560 1580
<ul><li>50</li><li>55</li><li>60</li><li>65</li></ul>	GAGCAACCTC CGACCCAGGG GCGGGGCCA ACCTGCCACC GCTGTTTGGGC GCCCCAGTGG GCACCAGTGG CATCCTCCTG CATCCTCCTG CATCCTCTG ATTCTATGAC TCTTGCAGGT ATTCTATGAC GAAAACAA GCAAATTGAC TTGATTTATATACA GTATGATTTATATACA TCATTATGAC TTATATTACT TATATATACA CTCATTATGT CATATTGT CATATTTTT CTAATTTACT TATATTACT TATATTACT TATATTACT TATATTACT TATATTACT CATATTTTT ATTCATTGGT AGCCAAGAAG GTGATAAATT TTTCATTGGT AGCCAAGAAG TTTTGATTACT TTTTCATTGGT TTTGATTTGA	11   AGCTTCTAGT CTTCTCCAGC GCCACCTTCG GCCACCTTCG CCTGAGCCAG TTCATTCTCG AGGATTACT TGGATGTCCT TGGATGTCCT CTGAATCTGA GGAGTGATAG GCAGTGATTACT CCTATGACCC GCTGCTTCTC ACCTCTTACC GTGTGACACA ATACTATATA TCATATCAT TGATACTAT TGATACTAT TGATACTAT TCATTACTC AAGGATGAAT CCATAATCTT CTCTATCTC AATTTATTC AAGGATGAAT CCTTATCTC AATTTATTAC CCTGTTTGACC AATATTTTT	21   ATCCAGACTC GGCGGCGGG GGAGTCCGGG CCCTTCCTGGG CCCTTCCTGGG GCGTGTCGCA GCAGCACATT CAATCTTTGT AGAAGATGAG TAGTTCCCTTC CAACACCAAG GAGGCAAAAG ATACATTAGT CAATACTTTTT ATACTTCATT ATACTTATAT TTCTTCATTA ATACTTAAAA ATTGGTAAAA TTCTTCATTA TCTTTCATTA TCTTTCATTA TCTTTCATTA TCTTTCATTA TCTTTCATTA TCTTTCATTA TCTTTCATTA TCTTTCATTA TCTTCATTA TCTTTCATTA TCTTTCATTA TCTTTCATTA TCTTCATTA TCTTCATTA TCAGACC CAAATCAGAC CCCACAC CCCACT	CAGCGCCGC CCAGCGAGGGC TTGCCCACCT CGAGCAGGGC TTGCCCACCT CGAGCAACATC GAGCACCGGG GCAACAACT GATGCTGTC AGCATGGTT CAGGTACGAA GGGAGGTGC GCCCTATCCA ACCTTAGAAT ACAAACCCAT TCCTCAATGT TTTCTCATGCGT CTTCATGGTT CATGTTTTC CTTTGAGTT TCTCATGAT TCATCATGTT ACATTTCATA TTTCTTTTC CTTTGGGTG TTTGGAGGCA ACCCTGTA ACCTTATAAA TTTCTTTTTC CTTTGGGTG CTTTGGGTG TTTGGGGTG TTTGGAGGCA ACCCTGTAC AGCTGCATGC	CCGGGCGCGG CCAAACTCTC ATGGCAACG GCCATCGTCA GTGACCGCCC CAGATCCATT ATTGGCTTGA GGCATGAAGT ATTGGGGTG ATTTGGGGTG ATTTGGTAAGAA ATTTGGTAATT GTGTAAAAT ATTGGGAAG GGAAGGGAA GGGAAGGGAA GGGAAGGGT ATAGAAACT TTTGGTTAAAAT ATTTGGTAAAAT ATTTGTTAAAAT CCTTTGCAC GCCCTTTTCA ACCTTATAT CCTTTGCAC GCCCTTTT AATCTTTCTC TCTGACCCAT TGTCCCCCA	ACCCCAACCC AACTTCTCC CGCCTTCTGC CGGCGTTCTGC CGGCGTTCTGC CGCCATGTA GCACAGCCT TCGTGCTTCT CGTTCAAC CTTCCTTCAC CTTCCAGCGG AACCCAAAAT CTATTTACT ATTTACAT GCATATTTACAT GCTCTTCAAC AATACTTAT GTATTTAAT AAATACTAT TGTATTTAAT TGATTTAAT TGATTTTATT TGATTTTATT TGATTTTATT TGATTTTATT TGATTTTATT TGATTTTATT TGATTTTATT TGATTTTATT TGATTTTATT TGATTTTATT TGATTTTATT TGATTTTATT TGATTTTATT TGATTTTATT TGATTTTATT TGATTTTATT TGATTTTATT TGATTTTATT TGATTTTATT TGATTTTATT TGATTTCTAA CATGACCAAA AGCACTCTTG GGTGTTGTAA	120 180 240 300 360 480 540 600 720 780 900 960 1020 1320 1380 1440 12560 1560 1560 1680 1740
<ul><li>50</li><li>55</li><li>60</li><li>65</li><li>70</li></ul>	GAGCAACCTC CGACCCAGGG GCGGGCCCA GCTGTTGGGC GCCCCAGTGG GCGCGCTGG CGAGGGGCTG TGACTCCTCTG CATCCTCTTG CATCCTCTG CATCCTCTTG CATCTCTATGAGT ATTCTATGAGT TGGAGAGAC TCTGCAGGT TGGATGAGC GGACATTGAG GTATGGTATT AAACATGGT TTGTATTACT TTATATTACT TATATTATTAC TTATTTTTTA TTCATTGGT TTGTTTTATT TTCATTGGT TTGCTTTGAT TTTGTTTGAT TTCATTGGT TTGCTTTGAT TTTGTTTGAT CCAAAAAA CAGCAAAAAA CTAATTTATTTTTTTTTT	11   AGCTTCTAGT CTTCTCCAGC CCCACCTTCG CCCACCTTCG CCCAGCCAC TCATTCTCG AGGATTTACT TGGATGTCCT CTGAATCTGA GAGGAGGATAT CCTATGACC CTGCTATTCC GCTGTTCTC ACCTCTTACC GTGTGACCAT ACAAACAAA TAATCTTATT GCTTCCCATT TAGGTATTATA TGATACTAGC GAAGATGTTT CCTTATCTC AAGGATGATT CCATTATCTC AAGGATGAT CCATATCTC CATTATCTC AAGTTGAC AAATTTTAT CCATTATTACTC CAATTATTAC CCTGTTGACC AAATTTTAT TGATTGACA AAATTTTTATTTAC CCAAATTTTTATTAC CCAAATTTTTTTT	21   ATCCAGACTC GGCGGCGCAG GGAGTCCGGG CCTATCCTGGG CCTATCCTGGG GCAGCACATT CAATCTTTGA TAGAAGATGA TAGTTAGCAC CAGTCAATGC CAGTCAATGC CAGTCAATGC CAGTCAATGC CAGTCAATGC CAGTCAATGC CAGTCAATGC TAGACACAAA AAGAACAATTAGG CAAACAACAA TTATCTTTT ATACTTAAAA ATTGGTATAT TCTTTCAATT ATAGGACTTA TGAATCTAAC TAAATCATAAC TAAATCATAAC TAAATCATAAC TTACATTAAAA ATTGGTATAT TCTTTCAATT ATAGCACTA CAAATCAGAAC TCCCACACA CAAATCAGACT TTTTAAAGCT	CAGCGCCGCC CGAGCAGGGC TTGCCACCT CGAGCGAGTC ATGGATCGGC CGACAACATC GAGCACCGT GATGGCTGTC AGCATCGTT AGCATCGGA GGAAGCAAC AGGAAGCAAC CAGGTACCAA CAGGTACCAA CAGGTACCAA CAGGTACCAA CACTTAGAAT AAAACCCAT TCCTCAAATAT ACTCTAAAT TCTCTTATC CTTTTTTTC CTTTTTTTT	CCGGGCGCGG CCGGCCCTT GCAAACTCTC ATGGCCAACG GCCATCGTCA GTGACCTCGC CAGATCCAGT ATTGGCGTG GCCATAGAA TTTGGTCAG GCCATTGTCA GGCAATAGAA TTTGGTCAGC CTACTTTGACAC TTTTGACAC GCGATCATA AGGAGGAAT ATAGGTAAAT AGGAGGAAG ATAGGTAAT CCTTTTGCCAC GCCTTTTCA AAGCCTTAT CCTTTTGCCAC GCCTTTTCA AAGCCTTAT TCTTGACCCAT TCTGACCCAT TCTGACCCAT GTTTCCCCCA GTTTTCATAT	ACCCCAACCC ACCTCTCTGC CGCGTTCTGC CGCGTTCTGC CGCGGTCTCA CCACTGCCCT AGGCCATGTA GCAAAGTCTT TGGTGGTTGG GTATGAAGA CTCTCTTCAC GTTCCTGTCC CTTCCAGCGG AACCGAAAAT ACTCAGTGCT ATTTTACCAT ATTTTACCAT ACTCTTCAA AAATACTATT GTATTTAATT ACATATGTAA AAGACCTAGC TATGTTTTATT TGTTTTTGTG TAGTTCTTAA CATGACCAAA AGGACCTATG CGGGGTTGTAA CATGACCAAA AGGACCTATG CGGGGTTGTAA CCCCTAAACT CCCCTAAACT CCCCTAAACT CCCCTAAACT	120 180 240 300 360 420 540 600 720 780 900 900 91080 1140 1260 1320 1320 1340 1500 1500 1680
<ul><li>50</li><li>55</li><li>60</li><li>65</li><li>70</li></ul>	GAGCAACCTC CGACCCAGGGGCCA ACCTGCCACC GCTGTTTGGGC GCCCCAGTGG GCCCCAGTGG CATCCTCCTG CATCCTCCTG CATCCTCTGCAGGT ATTCTATGAC TGGCTGGGCT ATTCTATGAC GAAAACAA GCAAATTACT TATATATACA TTGATTTACT TATATATACA CTCATTATGT CATATTGT CATATTGT CATATTGT CATATTGT TATATTACT TATATTACT TATATTACT TATATTACT TATATTACT TATATTACT AGCCAAGAAG GTGATAAATT TTTCATTTGT AGCCAAGAAG TTTGTATTTTA TTTTGTTTGA CACAACTTTTA CCATTTTTTA TTTTGTTTTG	11   AGCTTCTAGT CTTCTCAGC GCCACCTTCG GCCACCTTCG CCTGAGCCAG TTCATTCTCG AGGATTACT TGGATGTCCT CTGAATCTGA GGAGTGATAG GCAGTGATTACT CCTATGACCC GCTGCTATTT CCTATGACCC GTGTGACACA ATACTATCAT ACAAAACAAA	21   ATCCAGACTC GGCGGCGCG GGAGTCCGGG GCAGTCCCGG GCGTGTCCTGGG GCGTGTCCCA GCAGCACATT CAATCTTTGT AGAAGATGAG TAGTTCCTTC CAACACCAAG GAGGCAAAAG ATACTTTCT CAACACCAAG AGGCAAAAC TAATCTTTCT TAACATTACT TACATTATATTTT ATACTTAAAT TTCTTCATTA TCTTTCATTA TCTTTCATTA TCTTCATTA TAGACCTTG TGAATCTAAC AAATCAGAAC CAATTGAGT TTCCACCAC CCAATTGAGT TTTAAGCAT TTAATGTTT TTAACTTAAC	CAGCGCCGCC CGAGCAGGGC TTGCCCACCT CGAGCAGGGC TTGCCCACCT CGAGCAACATC GAGCACCGGG GCAACAACATC GAGCACCGTT GATGGCTGTC AGCATGGTACA GGGAGGTGCC GCCCTATCCA AGCATCCA ACCTTAGAAT ACAAAACCCAT TCCTCAATAT ACTCAAATGG TCTTATTAAA TATCTCTAAA TTTCTTTTC GCTTTGGGTG CTTCATGGTT ACATTTCATACAT TTTGGAGGCA ACCCTGCTAC AGCTGCATGC AGCTGCATGC AGCTGCATGC CTTATTCATA TTTTCATGCGTC CTTTATTCATA TTTTCATGCGTC CTTTATTCATA TTTTCATGCGTC CTTTTTCATCCTA AGCTGCATGC CTTATTCATA TTTTCATAC TTTTCCCA GCTGCATGC CTTTATTCCCA GCTCTGAACAA	CCGGGCGCGG CCAAACTCTC ATGGCAACG GCCATCGTCA GTGACCGCCC CAGATCCATGA GGCAATGAAGT ATTGGGGTG ATTGGGGTG ATTGGGGTG ATTGGGGTG ATTGGGTATAGAA TTTGGTAAAAT TTTGGTAAAAT ATTGGAAGA ATAGAAAA GCGAAGAAAA ATAGAAAAT GTCTTTAAA TCTTTTCTC CCCTTTTCA AACCTTATAT CCTTTGCCAC GCCTTTTCA ATTTTTCTG AATCTTTCTC AATCTTTCTC AATCTTTCTC AATCTTTCCAC GTTTTAATAT AGTGCAATTA AGTGCTAATT AGTGCTAATT AGTGCTAATT AGTGCTAATT AGTGCTAATT AGTGCTAATT AGTGCTAATT AGTGCTAAATT AGTGCTAAATT	ACCCCAACCC AACTTCTCC CGCCTTCTGC CGGCGTTCTGC CGGCGTTCTGC CGCGCTTCTGC CGCCATGTA GCAAAGTCTT TGGTGGTTGG CGATATTTCT TCGTTCAAG CTTCCTTCAC CTTCCAGCGG AACCGAAAAT GTAATCTGAA AATACTAT GTATTTAACTAT TGATTTTACAT TGATTTAAT TACATATTTTACT TAGTTTTATT TTATTTATT TAGTTTTTTTT TTGTTTTTTTT	120 180 300 360 480 640 720 780 900 900 1020 1140 1260 1380 1440 1556 1620 1680 1740 1800
<ul><li>50</li><li>55</li><li>60</li><li>65</li><li>70</li><li>75</li></ul>	GAGCAACCTC CGACCCAGGG GCGGGGCCCA ACCTGCCACC GCTGTTGGGC GCCCCAGTGG CGAGGGGCTG TGACTCCTCTG CATCCTCCTG CATCCTCTG CATCCTCTG CATCCAGGT ATTCTATGAC TCGGAGAAAACA GGACATTGAG GTATGGTATT AAACATGGCT TTGATTACT TTATATATAC CTATTTATTAC TTATTTTTTA CTTATTTTTTA ACCAGACATTGAG GTAGAGAAAAAC TCATTTGT TGGTATATTTTTTA TTTCATTGT TGCCAAGAAA GTGATAAATT TTTGCTTTGA CACAACTTTA ACCTTTTTTTA ACCTTTTTTTTA ACCTTTTTTTT	11   AGCTTCTAGT CTTCTCCAGC CCCACCATTCG CCCAGCCAG CCACCATTCG AGGATTTACT TGGATGTCCT CTGAATCTGA GGAGTGATAC CTGAATCTGA GCAGTGATTAC CTGATTCTC ACCTCTTACC GCTGTTCTC ACCTCTTACC GTGTGACACA ATACTATCAT ACAAAACAAA	21   ATCCAGACTC GGCGGCGCAG GGAGTCCGGG CCTATCCTGGG CCTATCCTGGG CCTATCCTGGG CCTATCCTGG CCAGCACATT CAATCTTTGT AGAAGATGAG TAGTCAATGC CAGTCAATGC CAGTCAATGC CAACACAAG GAGGCAAAAG ATACATTAGG CAACACAACA	CAGCGCCGC CCAGCAGGGC TTGCCACCT CGAGCAGGC TTGCCACCT CGAGCAGCC CGACAACATC GAGCACCGGC GCCACCGTT GATGGCTGTC AGCATGGGAC GGCACACGTC GACACGGT CAGGAACATC GAGGAACATC GAGGAACATC GCCTATCCA GAGAAAATC ACCTTAGAAT AAAACCCAT TCCTCAATAG TCTTATAAA TTTCTTTAG CTTCATGGTG CATCGTTAT TTTGGAGGCA ATCCCTGAAC ATCCCTGTAC ATCCCTGTAC TCTTATCATA TTTGGAGGCA ATCCCTGTAC TCTTATCATA TTTGGAGGCA TCTTATCATA TTTGGAGGCA TCCCTGTAC TCTTATCATA TGTTTCCCA GCTTTACACA GCTGTACCA GCTGTACCA GCTGTACACA GCTGTACACA	CCGGGCGCGG CCAACCTCT ATGCCAACG GCCATCGTCA ATGGCCACG GCATCGTCA GTGACCTTGA ATTGGCGTG GCAATAGAA ATTTGGTCAGG CTACTTTGCT AAACCTGCAC TTTTGAACA TTTGGGTATT GTGTAAAAT AGGAGGAAA AGGAGGAA ATAGGTAAT CCTTTTCAA ATTTGCCAC GCCTTTTA ATCTTATT CTCTTACACA TTTTGCCAC GCCTTTTCA AAGCCTTAT TCTTGCCAC GCCTTTTCA AGTCACATTT AATCTTTCTG TCTGACCCAT TGTTCCCCCA GTTTTATATC AGTGTAATT AGTTAATT AGTTAATT AGTTAATT AGTTAATT AGTTAATT AGTTAATT AGTTAATT AGTTAATT AGTTAATT AGTGTAATTA	ACCCCAACCC ACCTCTCTCC CGCCTTCTGC CGCGTTCTGC CGCGTTCTGC CGCGTGCAT GCACTGCCCT AGGCCATGTA GCAAGTCTT TGGTGGTTGG GTATGAAGT CGATATTCT TCGTTCAACG ACCGACAAT ACTCATCTCAC GTTCCTGCC CTTCCAGCG AACCGAAAAT GTAATCTGAA ACTCAGTGCT ATTTTACCAT GTATTTAATT ACATATGTATT ACATATTATT TTGTTTTTTT TTGTTTTTTT TTGTTTTTTT TGGTTCTTAA CATGCCAAA AGCACTCTG GGGGTTTTTAA CATGACCAAA AGCACTCTG GGGGTTTGTAA CCCCTAAACT TCATGCGTT TCTTCTGCAGT TCTTTCTGCAGT	120 180 300 360 480 660 720 780 900 900 1020 1140 1260 1320 1440 1550 1620 1680 1740 1800 1800 1980 1980
<ul><li>50</li><li>55</li><li>60</li><li>65</li><li>70</li></ul>	GAGCAACCTC CGACCAGCC GCTGTTGGGC GCCCCAGTGG GCGGGGGCTG TGACTCCTTG CATCCTCTG CATCCTCTG CATCCTCTG ATTCTATGAC TCTTGCAGGT ATTCTATGAC TGGTGGGCT GGACAGAGAC TCTTGCAGGT ATTCTATGAC GAAAAACA GAAAATAC GGACATTAAC TTGATTATTAC TTATATATAGA CTCATATTGT TTATATTATC TTTATTTTGT TTTCATTGG TTTGCTTTGA ACCAAGAAG GTGAAAATT TTTGCTTTGA ACCATTTTGT TTTGCTTTGT TATATATTTCC GATAATCTCC GATAATCTCC GATAATCTCC GATAATCTCC GATAATCTGT	11   AGCTTCTAGT CTTCTCCAGC GCCACCTTCG GCCACCTTCG CCTGAGCCAG TTCATTCTCG AGGATTTACT TGGATGTCTC CTGAATCTGA GAGGAGGAGC GATGATATC CCTGATTCTC ACCTCTTACC GCTGCTATTT CCTATGACCC GTGTGACACA ACAAACAAA TAATCTATCAT TAGTTACTAT TAGTTACTAT TAGTTACTAC AAGGATGTTT CCTTACTCC AAGGATGTTT CTCATTTACTC AAGGATGATT CCTTATCTC AATTATTT CTCTTATCTC AAATTTTT CTCTTATCTC AAATTTTT TCATTTACTC AAATTTTT TCATTTACTC AAATTTTTT TCATTTACTC AAATTTTTT TCATTTACTC AAATTTTTT TCATTTACTC AAATTTTTT TCATTTACTC CAAATTTTTT TTGATTTACAC CCTGTTGACC AAATTTTGT TTGATTGAAT TCCCCATTCC TAATAAGGTG TGACAAATAT ATCTGCCAAA	21   ATCCAGACTC GGCGGCGCAG GGAGTCCGGG CCTATGCCGG GCAGGCGCCC CCTTCCTGGG CCTATGCCGG GCAGCACATT CAATCTTTGT AGAGAATGAC TAGTTGCAC CAGTCAATGC CAACACAAG GAGCAAAAG TAACATTAGG TAACATTAGG TAACATTAGT TACATGTTT ATACTTAAAA ATTGGTATAT TCTTTCAATT ATAGACTAG TAAATCATT TTAAATCATT ATAGAACTAG TAAATCATT TTTCAATT TTCATTATT TTCTTCAATT TTCAATT TTCAATT TTTAAAT TTTTCAATT TTTAAAT TTTTCAATT TTTAAATTTT TTTAAGTAT TTTTAAGCTA TTTTAAGCTA TTTAAACTAT TTGTTTTT TTGTTTTT TTGTTTT TTGTTTTT TTGTTTT TTGTTTTT TTGTTTTT TTGTTTTTT	CAGCGCCGCC CGAGCAGGGC TTGCCCACCT CGAGCAGGGC TTGCCCACCT CGAGCAGTC GAGCACATC GAGCACCGT GATGCATCGT GATGCTGTC CAGCTACCAC GCCACCGTT ATGCTTACCA GCGAGAGCACC GCCCTATCCA GCAGAGCACC GCCCTATCCA ACCTTAGAAT AAAAACCCAT TCCTCAATAT ACTCAATAGA TTTCTTTTTC GCTTTGGGTC CATCGTTATTAAA TTTCTTTTTC CTTTAGGGT CATCGTTATTAAA TTTGTTTTTC CTTCATCATC ACCTTCATCAT ACATTCCTA ACATTCCTA ACCTTCATC ACCTTCATC ACCTTCATC ACCTTCATC ACCTTCATC ACCTTCATC ACCTTCATC ACCTTCATC ACCTTCATC ACCTTCATC ACCTTCATC ACCTTCATC ACCTTCATC ACCTTCATCATA ACCTTCATACA ACCTTCATACA ACCTTCATACA ACCTTCATACC ACCTTCATACA ACCTTCATACC ACCTTCATACC ACCTTCATACC ACCTTCATACC ACCTTCATACC ACCTTCATACC ACCTTCATACC ACCTTCATACC ACCTTCATACC ACCTTCATACC ACCTTCATACC ACCTTCATACC ACCTTCATACC ACCTTCATACC ACCTTCATACC ACCTTCATACC ACCTTCATACC ACCTTCATACC ACCTTCATACC ACCTTCATACC ACCTTCATACC ACCTTCATACC ACCTTCATACC ACCTTCATACC ACCTTCATACC ACCTTCATACC ACCTTCATACC ACCTTCATACC ACCTTCATACC ACCTTCATACC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC ACCTTCATACAC	CCGGGCGCGG CCAAACTCTC ATGGCCAACG GCCATCGTCA GCGATCGTCA GTGACCACCC CAGATCCACT GCTACCACT ATTGGGCTTA ATTGGGGTTA ATTGGGCTTA ATTGGGTAT TTTGGTCACA TTTTGAAACA TTTTGAAACA TTTTGAAACA TTTTAAACACACACACACACACACACACACACAC	ACCCCAACCC ACCTCTCC CGCCTTCTCC CGCCTTCTCC CGCGCTTCTCC CGCGCTTCTCC CGCGCTTCTCC CGCGCTTCTA GCAAAGTCTT TCGTTGATTCT TCGTTCACGTTCCTCCTCCC CTTCCAGCG AACCCGAAAAT GTAATCTAT GCAAATATTAT TCGTTCACGTC ATTTTACCAT GCTCCTTAAA AAATACTATT TCTTTTACCAT TTATTTAT	120 180 240 300 360 420 660 720 780 900 900 1020 1140 1260 1320 1380 1440 1500 1560 1740 1680 1740 1860 1980 1980
<ul><li>50</li><li>55</li><li>60</li><li>65</li><li>70</li><li>75</li></ul>	GAGCAACCTC CGACCCAGGG GCGGGGCCA ACCTGCCACC GCTGTTTGGGC CGCCCAGTGG CGAGGGGCTG CATCCTCCTG CATCCTCTG CATCCTCTG ATTCTATGAC TCTTGCAGTT ATTCTATGAC GAAAGACTAC GGACATTGAG GTATGGTATT AAACATGGCT TTGTATTACT TATATATACA CTCATTATGAC TCATTATGAT CATATTGAT CATATTGAT CATATGAT TCATTAGGT AGCCAAGAAGA GTGATAAATA TTTCATTGGT AGCCAAGAAGT ACCATCTTTTA TTTCATTTGAT TTTCATTTGAT TTTCATTTGAT TATATTTTA TTTCATTTGAT ACCAACTTTTA ACCATCTTTTTA TTTATATCTTTCC GATAATCTTCC GATAATCTTCC GATAATCTTCC GATAATCTTCC ATATTTTTT	11   AGCTTCTAGT CTTCTCAGC GCCACCTTCG GCCACCTTCG CCTGAGCCAG TTCATTCTCG AGGATTTACT TGGATGTCCT TGGATGTCCT CTGAATCTGA GGAGTGATAG GCAGTGATATT CCTATGACCC GCTGCTTTACC GTGTATACAT ACAAACAAA TAATCTTATT TGCTTCCCATT TATGTATATA TGATACTAC CAGAGTGATAT CCATAATCTC AAGGATGAT TCATTTACTC AAGGATGAT TCATTTACTC AAGTATATT CCTTATTCC AATTTATTAC CCTGTTGACC AAATATTTT TCATTTACTC TCATTTACTC TCATTTACTC TCATTTACTC TCATTTACTC TTATTACTC TCTCCCAAT ACTTCTCCCAAT CTTCTCCCAAT ACTTCTCCCAAT ACTTCTCCCAAT ACTTCTCCCAAT ACTTCTCCAAT ACTTCTCCCAAT ACTTCTCCAAT ACTTCTCCAAT ACTTCTCCAAT ACTTCTCCAAT ACTTCTCCAAT ACTTCTCCAAT ACTTCTCCAAT ACTTCTCCAAT ACTTCTCCAAT ACTTCTCCCAAT ACTTCTCCCAAT ACTTCTCCCAAT ACTTCTCCCAAT ACTTCTCCCAAT ACTTCTCCCAAT ACTTCTCCCAAT ACTTCTCCCAAT ACTTCTCTCCAAT ACTTCTCCAAT ACTTCTCCCAAT ACTTCTCTCCAAT ACTTCTCTCCAAT ACTTCTCTCCAAT ACTTCTCTCAA	21   ATCCAGACTC GGCGGCGGG GGAGTCCGGG GCAGTCCCAGG CCTATTCCTGGG GCGTGTCGCA GCAGCACATT CAATCTTTGT AGAAGATGAG TAGTTCCCTTCT CAACACCAG GAGGCAATAT CATCATTCT CAACACCAAG GAGGCAAAAC ATACTTATT TACATCTTCT TACATCTTCTT ATACTTCATT ATACTTATAT TCTTCATTA TCTTCATTA TCTTCATTA TCTTCATTA TCAGACCACA CCAATTGAGT TTAAGCACT TTAACTACAC CCAATTGAGT TTTAAGCTA TTAATTGTTT TGGTCTGTTT TCGTCTGTTT TCGTCTGTTT TCGTCTGTTA TTGGAGATAAT ACTCTCATTC	CAGCGCCGCC CGAGCAGGGC TTGCCCACCT CGAGCAGGGC TTGCCCACCT CGAGCAACATC GAGCACCGGG GCACACCGTT GATGGCTCT GATGGCTGTAT CAGGTACGAA GGGAGGTGCC GCCCTATCCA ACCTTAGAAT ACACTAGAAT TCCTCAATAT ACTCAATAT ACTCAATAT TTTCATGGGTG CTTCATGAT TTTGGAGGCA ACCCTGTTCATCAA TTTCATCATA TTTCATCATA TTTGGAGGCA ACCCTGTACCA ACCCTGTACCA GCTGCACGT CATGTATTCATA TTTGCAGGTG CTTATTCATA TTTGCAGGCG CTTATTCATA TTTGCAGGCA ACCCTGTAC AGCTGCATGC CTTGAACAA GCTGTAACAA GCTGTAACAA GCTGTAACAA GCTGTAACAA GCTGTAACAA GCTGTAACAA GCTGTAACAA TTTGAACATG	CCGGGCGCGG CCAAACTCTC ATGGCAACG GCCATCGTCA GTGACCGCCC GGCATCGTCA GTGACCGCCC CAGATCCAGT ATTGGGGTG ATTGGGGTG ATTGGGGTG ATTGGGGTG ATTGGGTAT ATTGGTCAGG CTACTTTGCT AAACCTGAAC ATTTGGTAAAAT ATGGGAAGGGAA	ACCCCAACCC AACTTCTCC CGCCTTCTGC CGCGTTCTGC CGGGGCTGCA GCACTGCCCT AGGCCATGTA GCAAAGTCTT TCGTGGTTGTG GTATGAAGTG CTCCTTCAC GTTCCAGCGG AACCGAAAAT GTAATCTGAA ACTACTGAT GCTCCTTAAA AAATACTATT GTATTTAACTAT GTATTTAACTAT TCGTTCTAGC TAGACTAGC CAGGGGTT TCATGCGTT TCATGCGTT TCATGCGTT TCATGCGTT TCATGCGTT TCATGCGTT TCATGCGTT TCATGCGTT TCATGCGTT TCATGCGTT TCATGCGTT TCATGCGTT TCATGCGTT TCATGCGTT TCATGCGTT TCATGCGTT TCATGCGTT TCATGCGTT TCATGCGTT TCATGCGTT AGGTAGTGT ATGTAGTTC AGGTAGTGT ATGTAGTGTC ATGTAGTTC AGGTAGTGT ATGTAGTGTC ATGTAGTGTC ATGTAGTGTC ATGTAGTGTC ATGTAGTGTC ATGTAGTAGTCT CTTCTACC AGGTAGTGTC ATGTAGTGTC ATGTAGTGTC ATGTAGTGTC ATGTAGTGTC ATGTAGTGTC ATGTAGTGTC ATGTAGTAGTCT CAGGCATT ATGTAGTGTC ATGTAGTAGTAGTAGTAGTAGTAGTAGTAGTAGTAGTAGT	120 180 240 300 360 420 540 600 720 780 960 1080 1140 1200 1320 1320 1320 1560 1560 1680 1740 1860 1920 1980 2040 2100
<ul><li>50</li><li>55</li><li>60</li><li>65</li><li>70</li><li>75</li></ul>	GAGCAACCTC CGACCAGCC GCTGTTGGGC GCCCCAGTGG GCGGGGGCTG TGACTCCTTG CATCCTCTG CATCCTCTG CATCCTCTG ATTCTATGAC TCTTGCAGGT ATTCTATGAC TGGCTGGGCT GGACAGCAGC GGACATTAT AAACATGGCT TTGTATTACT TATATATAGA CTAATTATTT TTTATTTGC TTTGCTTTGT TTTCTTTGT TTGCTTTGT TTTCTTTGT TTTGCTTTGT TTTTTTTT	11   AGCTTCTAGT CTTCTCCAGC GCCACCTTCG GCCACCTTCG CCTGAGCCAG TTCATTCTCG AGGATTTACT TGGATGTCTC CTGAATCTGA GAGGAGAGA CTGCTATTT CCTATGACCC GCTGCTATTT CCTATGACCA ACATACAT ACATACAT ACATACAT TAGTTACTT TAGTTACTAT TAGTTACTAT TAGTTACTAT TAGTTACTAT AAGGATGATT TCATTTACTC AAGGATGATT CCCTATCCC AATTTATTC CTCTATCTC CATATTTC TCATTTACTC AAATATTATT TCATTTACTC AAATATTATT TCATTTACTC CATATTTC CCTGTTGACC AAATATTTGT TGATTAGAT TCCCCATTCC TAATAAGGTG TGACAAATAT ATCTGCCAAA AGTTTATATT CAGCTGCCTAAA AGTTTATATT CAGCTGCCTT TCACTGCCTT TCACTGCCTT TCACTGCCTT TCACTGCCTT TCACTGCCTT TCACTGCCTT TCACTGCCTT TTCACTGCCTT TCACTGCCTT CACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCCT TCACTGCT TCACTGCCT TCACTGCCT TCACTGCT TCACTGCT TCACTGCCT TCACTGCT TCACTGCCT TCACTGCT TCACTGCT TCACTGCT TCACTGCT TCACTGCT TCACTGCT TCACTGCT TC	21   ATCCAGACTC GGCGGCGCAG GGAGTCCGGG CGCGGGGCGC CCTTCCTGGG GCAGCACATT CAATCTTTGT AGAGAATGA TAGTTGCAC CAGTCAATGC CAGTCAATGC CAACACAAG GAGCAAAAG GAGCAAAAG TAACATTAGG TAACATTAGG TAACATTAGT TACATTATAT ATTCTTCAATT ATTCTTCAATT ATTCTTCAATT ATAGACTAG TAACATTAGAT TTCTTCAATT TTCTTCAATT TTCTTCAATT TTCTTCAATT TTTAAGCAT TTTAAGCT TTTAAGCT TTTAAGCT TTTAAGCT TTTAAGCT TTTAAGCT TTTAAGCT TTTAAGCT TTGAGTATT TTGTCTCTTGT TTGAGTATT TTGTCTCTTT TTGAGATAAT AGCACACTC AGACACTGAT AGACACTGAT AGACACTGAT AGACACTGAT AGACACTGAT AGACACTGAT AGACACTGAT AGACACTGAT AGACACTGAT AGACACTGAT AGACACTGAT AGACACTGAT AGACACTGAT AGACACTGAT AGACACTGAT ACCCCCCTCTCT	CAGCGCCGCC CGAGCAGGGC TTGCCCACCT CGAGCAGGGC TTGCCCACCT CGAGCAGTC GAGCACTC GAGCACCGT GAGCACCGT GATGCTGTAC CAGCTACGAC GGCCACCGTT CATGCTACCA GGCAACCAC GCCCTATCCA GCAGAAAATCA ACCTTAGAAT ACAAATCCAT TCCTCAATAT ACTCAAATGG CTTCATTAAAA TTTCTTTTTC CCTTTGGGTC CATCGTTACTACA AGCTGCATC AGCTGCATC TTTGCACATC AGCTGCATC CTTCATTACAA TTTCTTTTC CTTTTGCCA TTTGGGCC CTTATTCATA ACTCCTGTAC AGCTGCATGC CTTATTCCAA AGCTGCATGC CTTATTCCAA AGCTGCATGC CTTATTCCAA GCTGTAACAA GCTGTAACA GCTGTAACA GCTGTAACC GAAGTCACTG GAAGTCACTG ACCAGTCTAT ACACTG GAAGTCACTG ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACCAGTCTAT ACC	CCGGGCGCGG CCAAACTCTC ATGGCCAACG GCCATCGTCA GTGACCGCCC CAGATCCATG GCATGAAGA ATTGGGCTAA ATTGGGGTAA ATTTGGGTAAA TTTGGTCAAGA CTACTTTGAAACA TTTGGGTATT GTGTTAAAAT AGGAGGAAG ATTAGACAGTA ATTGGCAC CCCTTATAT CCTTATAT CCTTATAT CCTTATAT TCTTAAACT TCTGCAC GCCCTTTTC AAGCCCTTAT TCTGACCCA GTTTTATAT ATTTCCCCA GTTTTATAT ATTTCCCCA GTTTTATAT ATTTCCCCA GTTTTATAT ATTTCCCCA GTTTTATAT ATTTCCCCA GTTTTATAT AGTGCACTA AACATTTA AGTGCTAAAA CAGTTAGAAA CAGTTAGAAG AACTATGCCT AACAAAACCT TTCCACTGAA	ACCCCAACCC ACCTCTCTC CGCCTTCTGC CGCGTTCTGC CGCGTTCTGC CGCGTTCTGC CGCGTGCCT AGGCCATGTA GCAAAGTCTT TCGTTGGTGTTGG GTATGAAGTG CTCTTCAC GTTCCTGTCC CTTCCAGCG AACCGAAAAT GTAATCTAT GCACAGAAAT GTAATCTAA ACTCAGTGCT ATTTTACCAT GCTCCTTAAA AAATACTATT TCTTTTATT TCATTTTATT TCATTTTTTTT	120 180 240 300 360 420 660 720 780 900 900 1020 1140 1260 1320 1380 1440 1500 1560 1740 1680 1740 1860 1980 1980
50 55 60 65 70 75	GAGCAACCTC CGACCCAGG GCGGGGCCC ACCTGCCACC GCTGTTGGGC CGAGGGGCTG CACCTCCTG CATCCCTG CATCCTGC CTTGGAAGAC TCTTGCAGGT ATTCTATGAC TGACTGCTG GGAAAACA GAAAGACTAC GGACATTGAG GTATGGTATT AAACATGGC TTGAATTACT TATATATATG CCATATTGAT CATATTTGT TCATTATGT TTTCATTGT TATATCTTG TATATCTTG TATATCTTGA CACAACTTTA ACCTTTTTTTTTT	11   AGCTTCTAGT CTTCTCCAGC GCCACCTTCG GCCACCTTCG CCTGAGCCAG TTCATTCTCG AGGATTTACT TGGATGTCCT TGGATGTGCT CTGAATCTGA GATGATGCC GCTGCTTCTC ACCTCTTACC GTGTGACACA ATACTATAT ACAAAACAAA	21   ATCCAGACTC GGCGGCGGG GGAGTCCGGG GCGGGCGCC CCTTCCTGGG GCGTGTCCCA GCAGCACATT AGAACATTTGT AGAAGATGA TAGTTGCAC CAGTCATTCT CAACCACAG GAGGCAAAAG ATATCTTCT CAACACAAG GAGGCAAAAG ATATCTTCT TAACATTAT TACATGTTTT ACATGTTTT ACATGTTAT TCTTCATTA TCTTCATTA TCTTCATTA TCTTCATTA TCTTCATTA TCTTCATTA TCTTCATTA TCTTCATTA TCTTCATTA TCTCACACA CCAATTGAGT TTTAAGCTTT TCTCTCTGTA TCGTCTCTTT TCTCTCTGTTA TCGTCTCTTT AGACACTGAA ACCCTCATTC AGACACTGAA ACCCTCCTCTC CAGTGCCTTC CAGTGCCTTC CAGTGCCTTC	CAGCGCCGCC CGAGCAGGGC TTGCCCACCT CGAGCAGGGC TTGCCCACCT CGAGCACTC GAGCACCGGG GCACACCGTT GATGGCTCT GATGGCTGTA CAGGTACGAA GGGAGGTGC GCCCTATCCA ACCTTAGAAT ACCTTAGAAT ACCTTAGAAT ACTCAATAT ACTCAATAT TTTCTTTTC GCTTCTAGAT TTTGGAGGC CTTATTCATA TTTGGAGGC CTTCATCCAA ATCCTTATCATA TTTGGAGGC CTTATTCATA TTTGGAGGC CTTATTCATA TTTGGAGGC AGCTGCATGC CTTATTCATA TTTGTATCCTA ACTTTATCATA TGGTTAT CTTTATCCTA GCTGCATGC CTTAATCAT GGTTGAACAA GCTGTAACAA TTTGAACAT GAAGCTCAT CTCTCTCTCAC	CCGGGCGCGG CCAAACTCTC ATGGCAACGCC GCAATCGTCA GTGACCGCCC CAGATCCATGA ATTGGGGTGA ATTGGGGTGA ATTGGGGTGA ATTGGGGTGA ATTGGGTAACG CTACTTTGCT AAACCTGCAC TGTTGAAACA TTTGGTAAAAC TTTGGTAAAAT ATTGGGAACA TTTGGTAAAAT GCGAATAGAAGT ATTAGACAGTA ATTGCCAC GCCTTTTCA AAGCCTTATA GCCTACTTTCA ATTCTTTCTG ATCTTTAAAT TCTTTCCCCC GCTTTTCA AGGCCTTATA AGGCCTTAT AATCTTTCTG AGGCCAT TGTCCCCCA GTTTTAATT AGTGCTACATTA AGTGCTAACA AGTCACTTAA AGTCACTTAA AGTCACTTAA AGTCACTTAA AGTCACTTAA AGTCACTTAA AGTCACTTAA AGTCACTTAA AGTCACTTAA AGTCACTTAA AGTCACTTAA AGTCACTTAA AGTCACTTAA AGTCACTTAA AGTCACTTAA AGTCACTTAA AGTCACTTAA AGTCACTTAA AGTCACTTAA CAGTTAGAAG AACTATGCCT ATTCACCCATTTT TCCACTGAA CAGTCTATTT	ACCCCAACCC ACCTCTCTC CGCCTTCTGC CGCGTTCTGC CGCGTTCTGC CGCGTTCTGC CGCGTGCCT AGGCCATGTA GCAAAGTCTT TCGTTGATGT CGTTCAAGT CTCCTTCAC GTTCCTGTCC CTTCCAGCGG AACCGAAAAT GTAATCTGAA ACTCAGTGCT ACTCAGTGCT TCGTTCAAG ACCGAAAAT GTAATCTAAT GTATTTAATT ACATATGTAA AAAACTTAT TTGTTTTTTTG TAGTTTCTAA CATGACCAAA AGCACTCTTG GGTGTTGTAA CCCCTAAACTTAT TCATGCGTTT TCATGCGTTT TCATGCGTTT TCATGCGTTT TCATGCGTTT TCATGCGTTT TCATGCGTTT TCATGCGTTT TCATGCGTTT TCATGCGTTT TCATGCGTTT CATGCGTTT ACACACCTAC CCACACCTAC CCACACCTAC CCACTGAACC	120 180 240 300 360 420 780 960 900 960 1080 1140 1260 1320 1320 1440 1500 1560 1740 1860 1920 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980
<ul><li>50</li><li>55</li><li>60</li><li>65</li><li>70</li><li>75</li></ul>	GAGCAACCTC CGACCCAGGG GCGGGGCCCA ACCTGCCACC GCTGTTGGGC GCCCCAGTGG CGCCCCAGTGG CGCCCCAGTGG CATCCTCCTG CATCCTCCTG CATCCTCTG CATCCTCTG CATCATCAGAGA TCTTGAAGAC TCTGCAGGT ATACATTAGA GTATGATTACT TATATTATACT TATATTATTACT TATATTATTA CTAATTTACT TATATTTTTA ACCTACTTTTTA ACCTACTTTTTA ACCTTTTTTA ACCTTTTTTA ACCTTTTTTA ACCTTTTTTTA ACCTTTTTTTA ACCTTTTTTTA ACCTTTTTTTA ACCTTTTTTTA ACCTTTTTTTT	11   AGCTTCTAGT CTTCTCCAGC CCTGAGCCAG CCCACCTTCG CCTGAGCCAG TCCATTCTC TGGATGTCCT TGGATGTCCT TGGATGTAC GAGGTGATA GATGAGGTGC CTGGCTATTT CCTATGACC GCTGCTATTC ACCTCTTACC GTGTGACACA ATACTATCAT ACAAACAAA TAATCTATT TGATCATC AGGATGATT TCATTACTC AAGGATGATT CCATATCC AAGTATTT CCTATCTC AATATTATT CCTTTACC AATATTATT TCTCTATCTC AATATTATT TCTCTTTACC CTGTTGACC AATATTTTT TCTTATTTAC CCTGTTGAC TCAATAAGGTG TTGACTAC TGACAAATAT ACTTGCCAAA ACTTTATTAC CAGCTGGCTG TCACTGCCTT CAGCTGGCTT CAGTTGCCTT CAGCTGGCTT CACTTCCC TCACTTCCC TTCACTTCC TTCACTTCC TTCACTTCC TTCACTTCC TTCACTTCC TTCACTTCC TTCACTTCC TTCACTTCC TTCACTTCC TTCACTTCC TTCACTTCC TTCACTTCC TTCACTTCC TTCACTTCC TCACTTCC CACTTC TCACTTC TCACTTC TCACTTC TCACTTC TCACTTC TCACTTC TCACTTC TCACTTC TCACTTC TCACTTC TCACTTC TCACTTC TCACTTC TCACTTC TCACTTC TCACTTC TCACTTC TCACTTC TCACTTC TCACTTC TCACTTC TCACTTC TCACTTC TCACTTC TCACTTC TCACTTC TCACTTC TCACTTC TCACTTC TCACTTC TCACTTC TCACTT TCACTTC TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACTT TCACT TCACT TCACT TCACT TCACT TCACT TCACT TCACT TCACT TCACT TCACT TCACT TCACT	21   ATCCAGACTC GGCGGCGCG GGAGTCCGGG GCAGGCCCC CCTTCCTGGG CCTATGCCGG GCGTGTCGCA GCAGCACATT CAATCTTTGT AGAAGATGAG TAGTCATCC CAGCACACA GAGGCAAAAG GAGGCAAAAG ATACTTTCT CAACACAAG AAGCAAACA TAATCTTCTT ATACTTAAAA ATTGGTATAT TCTTCATTA TCTTCATTA TCTTCAATT ATACTTCATTA TCAATCTTCATT ATACACTTG TGAATCATAC AAATCAGAC CCAATTGAGT TTTAAGCACTTG TGAATCTTAAC TTAATCTTCTT TGAATCTTAAC TTAATCTTCTT TGAATCTAAC AAATCAGAC TCCCACCA CCAATTGAGT TTTAAGCTA TCCTCTGTT TCGCTCTGTA TGGCTCTCTT AGGCACTCA AGCACTGAA TCCTCTCTC AGGCACTC AGGCCTTC AGGTGCCTTC AGTGGCCTC ATGTGGCTTC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGCTCC ATGTGGCTCC ATGTGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGGCTCC ATGTGCTCC ATGTGCCTCC ATGTTCC ATGTGCCTCC ATGTGCCTCC ATGTGCCTCC ATGTGCCTCC ATGTGCCTCC ATGTGC	CAGCGCCGCC CGAGCAGGGC TTGCCCACCT CGAGCAGGGC TTGCCCACCT CGAGCACATC GAGCACCGGG GCAACAACATC GAGCACCGTT GATGGCTGTC AGCATGGTAC GGCACCGTT CAAGTACGAC GGCACCGTT CAAGTACGAC GGCAACATCA GGAAAAATCA AACACTAT AAAAACCCAT TCCTCAAATTG CTTCATGGGT CTTCATGGGT CTTCATGGGT CTTCATGGT CTTCATGGT CTTCATGAT TCCTGATTT ACATTTCATA TTTCCTTATA TTTCTTTTC CTTCATGGT CTTTTCATGGT CTTTTTCATA TTTGGAGGCA ACCTTATTCATA TTTGAACATG GTTTTTCCTGACGT CTTATTCATA TGTTTTCCTGACGT CTTATTCATA TTTTCCTGACGA GCTGCATGC CTTATTCATA TCTTTTCCTGCT GACAGCTCAT CCCGGTCAT CCTCTCTCC CTGCCTTCCT CTGCCTTCCT CTGCCTTCCT CTGCCTTCCT CTGCCTTCCT CTGCCTTCCT CTGCCTTCCT CTGCCTTCCT CTGCCTTCCT CTGCCTTCCT CTGCCTTCCT CTGCCTTCCT CTGCCTTCCT CTGCCTTCCT CTGCCTTCCT CTGCCTTCCT CTGCCTTCCT CTGCCTTCCT CTGCCTTCCT CTGCCTTCCT CTGCCTTCCT CTGCCTTCCT CTGCCACC CTGCCTTCCT CTGCCTCCT CTGCCACC CTGCCTTCCT CTGCCACC CTGCCTTCCT CTGCCTCCT CTGCCACC CTGCCTTCCT CTGCCACC CTGCCTTCCT CTGCCTCCT CTGCCTTCCT CTGCCACC CTGCCTTCCT CTGCCACC CTGCCACC CTGCCTCCT CTGCCACC CTGCCTTCCT CTGCCACC CTGCCTTCCT CTGCCTCCT CTGCCACC CTGCCTTCCT CTGCCTTCCT CTGCCTTCCT CTGCCTTCCT CTGCCTTCCT CTGCCTTCCT CTGCCTTCCT CTGCCTTCCT CTGCCTTCCT CTGCCTTCCT CTGCCTTCCT CTGCCTTCCT CTGCCTTCCT CTGCCTTCCT CTGCCTTCCT CTGCCTTCCT CTGCCTTCCT CTGCCTTCCT CTGCCTTCCT CTGCCTTCCT CTGCCTTCCT CTGCCTTCCT CTGCCTTCCT CTGCCTTCCT CTCCTCTCTCT	CCGGGCGCGG CCGGCCTT GCAAACTCTC ATGGCCAACG GCCATCGTCA GTGACCGCCC CAGATCCATGA GGCATGAAGT ATTGGGGTG ATTTGGGGTG TTTGGTCATGA GCAATAGAA TTTGGTCATGA TTTGGTCATG TTTGGTATT GTGTTAAAAT AGGAGGAGA ATAGGAATAAA AGGAGGAGA TCTTTGCCAC GCCTTTTCA CCTTTCCAC GTTTTAATC AGTGAACT TGTTCCCCCA GTTTTATATC AGTGTAATT AGTGACTATT AGTGTAATT AGTGACCTTA AGTGTAATT CCTTTCCCCA GTTTTAATC AGTGTAATT CCTTTCCCCA GTTTTAATC AGTGTAATT CAGTGAACA TTTCCCCCA TTTTCCCCCA GTTTTAATC CAGTGAACA CAGTTAAACCT TCCCCCTACATTT AGTGTAATT CCTTCCCCCA CGTTTTAATC CGCTACTTAA CGGTAATTC CCTCCCCA CGTTTTAATC CGCTCCCA CGTTTTAATC CGCTCCCA CGTTTAACCCT ACAAAACCT TCCCCCTGAA CAGTCTATTT CCTCTCCCCCA CTCCTCCCCCC CTCCCTCC	ACCCCAACCC ACCTCTCTC CGCCTTCTGC CGCGTTCTGC CGCGTTCTGC CGCGTGCCT AGGCCATGTA GCAAAGTCTT TGGTGGTTGG GTATGAAGTG CGATATTTCT TCGTTCAAG CTTCCTTCAC CTTCCAGCGG AACCGAAAAT GTAATCTGAA ACTCAGTGCT ATTTTACCAT ACTCAGTGTAAA AAATACTAAT GTATTTAATT GTATTTAATT TGATTTTATT TGTATTTATT	120 180 300 360 480 540 600 720 840 900 1020 1140 1260 1320 1440 1550 1620 1620 1740 1800 1740 1800 1980 2040 2160 2220 22340
50 55 60 65 70 75	GAGCAACCTC CGACCCAGG GCGGGCCCA GCTGTTGGGC GCCCCAGTGG GCGCGCTG TGACTCCTCTG CATCCTCTG CATCCTCTG CATCCTCTG CATCCTCTG CATCCTCTG CATCTCATGAG TTGAAGAC TCTTGCAGGT ATTCTATGAC GGACATTGAG GTATGGTATT AAACATGGCT TTGATTACT TTATTATTAC TTATTTTTA TTTCATTGGT TATTTTTTA TTTCATTGGT TATCTTTGA TTTGCTTTGA CACAACTTTA ACCTTTTTG TATATCTTCC TATATCTTC TATATCTTCT TATATCTTCT TATATCTTCT TATATCTTCC TATATCTTCC TATATCTTCC TATATCTTCC TATATCTTCC TATATCTTCC TATATCTTCC TATATCTTCC AATATCTGC TCTTTTTCT TTTATTTTCT TTTATTTTCT TTTATTTTCT TTTATTTTCT TTTATTTTCT TTTATTTTCT TTTATTTTCT TTTATTTTCT TTTATTTTCT TTTATTTCT AAACCTACGC AATCCTT	11   AGCTTCTAGT CTTCTCCAGC GCCACCTTCG GCCACCTTCG CCTGAGCCAG TTCATTCTCG AGGATTTACT TGGATGTCCT TGGATGTGCT CTGAATCTGA GATGATGCC GCTGCTTCTC ACCTCTTACC GTGTGACACA ATACTATAT ACAAAACAAA	21   ATCCAGACTC GGCGGCGCAG GGAGTCCGGG GCGGGGCGC CCTTCCTGGG GCGTGTCCGG GCAGCACATT CAATCTTTG AGAGACACT TAGACGACAA GAGCACAAG GAGCACAAG GAGCACAAG GAGCACAAG GAGCACAAG GAGCACAAG GAGCACAAG ACACACAAG TAACATTAGG 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1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 1980 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55
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                                                                                                1020
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70
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         RABISHHISS CODRSCIECD VVNQTRSLRQ ETLABSTWQC PPCDEDWDKD LWEQTSTPFV WGTTHYSDNN SPASNIVTEH KNNLASGMRV PKSLPYVLPW KNNGNAQ
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WO 02/086443
Seq ID NO: 248 DNA sequence
Nucleic Acid Accession #: NM_003392
Coding sequence: 758..1855

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	TO COTO COOT	GCCCCGCGCA	CAGGATCCCA	GCGAAAATCA	GATTTCCTGG	TGAGGTTGCG	540
	TOCOTOGATT	ΔΔΑΣΣΤΈΤΑΔ	AAGAAACTGC	CTATATCTTG	CCATCAAAAA	ACTCACGGAG	600
	GAGAAGCGCA	GTCAATCAAC	AGTAAACTTA	AGAGACCCCC	GATGCTCCCC	TGGTTTAACT	660 720
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	CAATTCTTGG	CCTCTCTGCA	CCCAACTCCC	AGGACTTTCT	CAAGGACAGA	AGAAACTGTG	960
	CON COMOUNTS	CACCACCACA	<b>ずばいなびかないなず</b>	CGGAGAAGGC	GCGAAGACAG	GCATCAAAGA	1020
25	ATCCCACTAT	CAATTCCGAC	ATCGACGGTG	GAACTGCAGC	ACTGTGGATA	ACACCTCTGT	1080
	<b>ササササビにことな</b>	CTCATCCAGA	TAGGCAGCCG	CGAGACGGCC	TTCACATACG	CCGTGAGCGC	1140
	ACCR CCCCTTC	CTCAACCCCA	TCAGCCGGGC	GTGCCGCGAG	GGCGAGCTGT	CCACCTGCGG	1200
•	CTCCNGCCGC	accececece.	CCAAGGACCT	GCCGCGGGAC	TGGCTCTGGG	GCGGCTGCGG	1260
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<i>J J</i>	CCTGTGCNAC	AAGACGTCGG	AGGGCATGGA	TGGCTGCGAG	CTCATGTGCT	GCGGCCGTGG	1740
	CEN CONCORC	TTCAACACCC	TOCAGACGGA	GCGCTGCCAC	TGCAAGTTCC	ACTGGTGCTG	1800
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45	CIGIGIGGA	TOCONTOCOT	ACCEMENTS	TGAAAGAGGG	TGGTAGAAAT	CTATTCACAA	2280
73	<b>サイベル スペヤヤペヤ</b>	ATCACCAAAA	TGAGTTGTAA	ATTCTCTGGT	GCAAGATAAA	AGGTCTTGGG	2340
	ממממממממ	DAAACAAAAC	AAACCTCCCT	TCCCCAGCAG	GGCTGCTAGC	TTGCTTTCTG	2400
	Characters	ስጥርስጥል ስ <b>ፐ</b> ፕፕ	ACAATGGAAG	GACAAGAATG	TCATATTCTC	AAGGAAAAA	2460
	COTATATO	<b>አጥርጥርጥር'ል</b> ጥጥ	CTCCTCAAAT	ATTCCATTTG	CAGACAGACC	GTCATATICI	2520
50	スカポルのクサイカオ	CABATTTCCC	CAGCAGGGAG	GAAAGTCCCC	AGAAATTAAA	AAATTTAAAA	2580
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	ATGAAATATC	CTGTATTTTC	TTAGGGATAC	CCAACCAACA	AGGTAATTGC	GTAGAAATAA GTGCCATTCA	2820
55	TACATGAATC	CCATTCACAG	ACCUMUTE TO THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION OF THE ACCUMUNTATION	CCAAGCAACA	TGAAATCCAC	CTTCCTCTTC	2880
33	ACACTGCACC	CTCTCTCTCTTT	CCTCCGTGTT	GTGATGTGAT	GCTGGCCACG	TTTCCAAACG	2940
	acaecrecoae	<b>ずににはずじじじて</b>	TTGGTTGTAG	GACAGGAAAT	GAAACATTAG	GAGCTCTGCT	3000
	TOGADADORG	ייידים ביים ביידית	AGGGATTTT	GTTTCCTAAA	ACTITIATI	TGAGGAGCAG	3060
	TO CONTRACT A	ጥርምምም ል ው	. ልሮልናልልሮሞፕና	GCTAATGGAA	TTCACAGAGG	TGTTGCAGCG	3120
60	ጥ አጥር አርጥር ጥጥ	TETTOTALDEA	GTTTAGATTA	TCCACTCATG	CTTCTCCTAT	TGTACTGCAG	3180
	OROMA COMMA	8 8 8 CTCTTCC	<b>・ へんごかごかる ごかか</b>	GAACAGTTGC	ATTTATAAGG	GGGGAAATGT	3240
	GGTTTAATGG	TGCCTGATAT	CTCAAAGTCT	TTTGTACATA	ACATATATAT	ATATATACAT	3300
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65	CTCTGGGGTT	ATCTCTCTGT	CTAGAGCATT	CCCTCTCCCCC	CCCCTCTGAT	AGTTGGGATT CATACCCTGA	3480
05			ע ענייט ע האייייייייייייייייייייייייייייייייייי	CANCETTONO	TTCTCACTC	CTGAAATGCG	3540
	TOTTCCCTTC	CONNECTED I	TTCTGAGGA	CTGCCTCACC	CCTTTGTCTC	CAACCTCCAT	3600
	<b>サマンナンサイン (************************************</b>	TTTTTGAGAGA	GGGCATTACT	TGTTCGTTAT	AGACATUGAC	; GTTAAGAGAT	3660
	ከግ <b>ፈፈፈፈ</b> ስግተዋል	CAGAAGCATC	AGCAATGTTT	CTCTTTTCTI	AGTTCATTC	GCAGAATGGA	3720
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	CACCCCATCA	እአጥአጥርያጥጥርጥ	' ጥልጥርጥር <b>ል</b> ርልር	TTACGTTGTT	TTAAAAGTTI	GGAAAGATAC	3900
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	CRTTCACATA	<b>የፈጥልጥርንፕልጥቤ</b>	CTTCTAGCCI	TTATTCTGT	CTTTTAATG	r ACATATTTCT	4320
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	AATGGAAGAT	AGAATATAA	ATAAAACGTT	ACTTGTAAA	AAAAAAA A		
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MAGSAMSSKF FLVALAIFFS FAQVVIEANS WWSLGMINPV QMSEVYIIGA QPLCSQLAGL
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TGTGTGTTCG ACAAAGAACC GGCGCTGCAA ACTGCTGCCC CTGGTGATGG CTGCCCCCCT
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Seq ID NO: 285 DNA sequence Nucleic Acid Accession #: Eos sequence Coding sequence: 1-1746

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## WO 02/086443

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## WO 02/086443

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Protein Accession #: Eos sequence

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ATATAACACT CAAAAAAATG TAAATCATAT TGTAGTATTC AATAGTTAAT AAAAACTCGA 1560 70 GAAATGTGTT GTTTCTG Seq ID NO: 321 Protein sequence: Protein Accession #: NP 005438.1 75 51 MAPFGRNLLK TRHKNRSPTK DMDSEEKBIV VWVCQEEKLV CGLTKRTTSA DVIQALLEEH 60 EATFGEKRPL LGKPSDYCII EKWRGSERVL PPLTRILKLW KAWGDEQPMM QFVLVKADAF LPVPLWRTAE AKLVQNTEKL WELSPANYMK TLPPDKQKRI VRKTFRKLAK IKQDTVSHDR 120 180 80 DNMETLVHLI ISQDHTIHQQ VKRMKELDLE IEKCEAKFHL DRVENDGENY VQDAYLMPSF 240 SEVEQUIDIO YEENOTLEDL SESDGIEQLE ERLKYYRILI DKLSAEIEKE VKSVCIDINE 300 DAEGRAASEL ESSNLESVKC DLEKSMKAGL KIHSHLSGIQ KEIKYSDSLL QMKAKEYELL AKEFNSLHIS NKDGCQLKEN RAKESEVPSS NGEIPPFTQR VFSNYTNDTD SDTGISSNHS 85 Seg ID NO: 322 DNA sequence

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Seq ID NO: 344 DNA sequence Nucleic Acid Accession #: NM_012072 Coding sequence: 149-2107

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WO 02/086443

Coding sequence: 1..927

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WO 02/086443

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WO 02/086443

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PCT/US02/12476

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		GCAAAATGCT					5400
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	Sec ID NO-	465 Protein	n semience				
80		ession #: 1					
	January 100					•	
	ļ	11	21	31	41	51	
				DIGITAL COL	III.I CAUTAMA	OUTERMENT	
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300

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TMGMVNIFNG DADLSGMTWS HGLSVSKVLH KAFVEVTEEG VEAAAATAVV VVELSSPSTN EEFCCNHPFL FFIRQNKTNS ILFYGRFSSP 5 Seq ID NO: 466 DNA sequence Nucleic Acid Accession #: NM_001910.1 Coding sequence: 50..1240 10 GGAGAGAAGA AAGGAGGGGG CAAGGGAGAA GCTGCTGGTC GGACTCACAA TGAAAACGCT CCTTCTTTTG CTGCTGGTGC TCCTGGAGCT GGGAGAGGCC CAAGGATCCC TTCACAGGGT GCCCCTCAGG AGGCATCCGT CCCTCAAGAA GAAGCTGCGG GCACGGAGCC AGCTCTCTGA 180 GTTCTGGAAA TCCCATAATT TGGACATGAT CCAGTTCACC GAGTCCTGCT CAATGGACCA 240 GAGTGCCAAG GAACCCCTCA TCAACTACTT GGATATGGAA TACTTCGGCA CTATCTCCAT TGGCTCCCCA CCACAGAACT TCACTGTCAT CTTCGACACT GGCTCCTCCA ACCTCTGGGT 15 300 CCCCTCTGTG TACTGCACTA GCCCAGCCTG CAAGACGCAC AGCAGGTTCC AGCCTTCCCA 420 GTCCAGCACA TACAGCCAGC CAGGTCAATC TTTCTCCATT CAGTATGGAA CCGGGAGCTT GTCCGGGATC ATTGGAGCCG ACCAAGTCTC TGTGGAAGGA CTAACCGTGG TTGGCCAGCA 480 540 20 GTTTGGAGAA AGTGTCACAG AGCCAGGCCA GACCTTTGTG GATGCAGAGT TTGATGGAAT 600 TCTGGGCCTG GGATACCCCT CCTTGGCTGT GGGAGGAGTG ACTCCAGTAT TTGACAACAT GATGGCTCAG AACCTGGTGG ACTTGCCGAT GTTTTCTGTC TACATGAGCA GTAACCCAGA 660 720 AGGTGGTGCG GGGAGCGAGC TGATTTTTGG AGGCTACGAC CACTCCCATT TCTCTGGGAG CCTGAATTGG GTCCCAGTCA CCAAGCAAGC TTACTGGCAG ATTGCACTGG ATAACATCCA 840 25 GGTGGGAGGC ACTGTTATGT TCTGCTCCGA GGGCTGCCAG GCCATTGTGG ACACAGGGAC 900 TTCCCTCATC ACTGGCCCTT CCGACAAGAT TAAGCAGCTG CAAAACGCCA TTGGGGCAGC 960 CCCCGTGGAT GGAGAATATG CTGTGGAGTG TGCCAACCTT AACGTCATGC CGGATGTCAC 1020 CTTCACCATT AACGGAGTCC CCTATACCCT CAGCCCAACT GCCTACACCC TACTGGACTT 1080 CGTGGATGGA ATGCAGTTCT GCAGCAGTGG CTTTCAAGGA CTTGACATCC ACCCTCCAGC TGGGCCCCTC TGGATCCTGG GGGATGTCTT CATTCGACAG TTTTACTCAG TCTTTGACCG 1140 30 1200 TGGGAATAAC CGTGTGGGAC TGGCCCCAGC AGTCCCCTAA GGAGGGGCCT TGTGTCTGTG 1260 CCTGCCTGTC TGACAGACCT TGAATATGTT AGGCTGGGGC ATTCTTTACA CCTACAAAAA GTTATTTTCC AGAGAATGTA GCTGTTTCCA GGGTTGCAAC TTGAATTAAG ACCAAACAGA ACATGAGAAT ACACACACA ACACACATAT ACACACACA ACACTTCACA CATACACACC 1320 1380 1440 35 ACTCCCACCA CCGTCATGAT GGAGGAATTA CGTTATACAT TCATATTTTG TATTGATTTT 1500 TGATTATGAA AATCAAAAAT TTTCACATTT GATTATGAAA ATCTCCAAAC ATATGCACAA GCAGAGATCA TGGTATAATA AATCCCTTTG CAACTCCACT CAGCCCTGAC AACCCATCCA CACACGGCCA GGCCTGTTTA TCTACACTGC TGCCCACTCC TCTCTCCAGC TCCACATGCT 1560 1620 1680 GTACCTGGAT CATTCTGAAG CAAATTCCGA GCATTACATC ATTTTGTCCA TAAATATTTC TAACATCCTT AAATATACAA TCGGAATTCA AGCATCTCCC ATTGTCCCAC AAATGTTTGG CTGTTTTTGT AGTTGGATTG TTTGTATTAG GATTCAAGCA AGGCCCATAT ATTGCATTTA 1740 40 1800 1860 TTTGAAATGT CTGTAAGTCT CTTTCCATCT ACAGAGTTTA GCACATTTGA ACGTTGCTGG 1920 TTGAAATCCC GAGGTGTCAT TTGACATGGT TCTCTGAACT TATCTTTCCT ATAAAATGGT 1980 AGTTAGATCT GGAGGTCTGA TTTTGTGGCA AAAATACTTC CTAGGTGGTG CTGGGTACTT 45 CTTGTTGCAT CCTGTCAGGA GGCAGATAAT GCTGGTGCCT CTCTATTGGT AATGTTAAGA 2100 CTGCTGGGTG GGTTTGGAGT TCTTGGCTTT AATCATTCAT TACAAAGTTC AGCATTTT Seg ID NO: 467 Protein sequence Protein Accession #: NP_001901.1 50 51 31 41 21 MKTLLLLLLV LLELGEAGGS LHRVPLRRHP SLKKKLRARS QLSEFWKSHN LDMIQFTESC SMDQSAKEPI INYLDMEYFG TISIGSPPQN FTVIFDTGSS NLWVPSVYCT SPACKTHSRF QPSQSSTYSQ PGQSFSIQYG TGSLSGIIGA DQVSVEGLTV VGQQFGESVT EPGQTFVDAE 55 180 FDGILGLGYP SLAVGGVTPV FDNMMAQNLV DLPMFSVYMS SNPEGGAGSE LIFGGYDHSH 240 FSGSLNWVPV TKQAYWQIAL DNIQVGGTVM FCSEGCQAIV DTGTSLITGP SDKIKQLQNA 300 IGAAPVDGBY AVECANLNVM PDVTFTINGV PYTLSPTAYT LLDFVDGMQF CSSGFQGLDI HPPAGPLWIL GDVFIRQFYS VFDRGNNRVG LAPAVP 60 Seg ID NO: 468 DNA seguence Nucleic Acid Accession #: NM_018058.1 Coding sequence: 319..1575 65 TACGCGCTGC GGGACCGGCA GGGGAACGCC ATCGGGGTCA CAGCCTGCGA CATCGACGGG GACGGCCGGG AGGAGATCTA CTTCCTCAAC ACCAATAATG CCTTCTCGGG GGTGGCCACG 120 TACACCGACA AGTTGTTCAA GTTCCGCAAT AACCGGTGGG AAGACATCCT GAGCGATGAG GTCAACGTGG CCCGTGGTGT GGCCAGCCTC TTTGCCGGAC GCTCTGTGGC CTGTGTGGAC 180 70 240 AGAAAGGGCT CTGGACGCTA CTCTATCTAC ATTGCCAATT ACGCCTACGG TAATGTGGGC 360 CCTGATGCCC TCATTGAAAT GGACCCTGAG GCCAGTGACC TCTCCCGGGG CATTCTGGCG CTCAGAGATO TEGCTECTGA GECTEGGETC AGCAAATATA CAGGGGGCCG AGGCGTCAGC GTGGGCCCCA TCCTCAGCAG CAGTGCCTCG GATATCTTCT GCGACAATGA GAATGGGCCT AACTTCCTTT TCCACAACCG GGGCGATGGC ACCTTTGTGG ACGCTGCGGC CAGTGCTGGT 420 480 75 GTGGACGACC CCCACCAGCA TGGGCGAGGT GTCGCCCTGG CTGACTTCAA CCGTGATGGC AAAGTGGACA TCGTCTATGG CAACTGGAAT GGCCCCCACC GCCTCTATCT GCAAATGAGC 600 660 ACCCATGGGA AGGTCCGCTT CCGGGACATC GCCTCACCCA AGTTCTCCAT GCCCTCCCCT GTCCGCACGG TCATCACCGC CGACTTTGAC AATGACCAGG AGCTGGAGAT CTTCTTCAAC AACATTGCCT ACCGCAGCTC CTCAGCCAAC CGCCTCTTCC GCGTCATCCG TAGAGAGCAC GGAGACCCCC TCATCGAGGA GCTCAATCCC GGCGACCCCT TGGAGCCTGA GGGCCGGGGC 780 80 840 900 ACAGGGGGTG TGGTGACCGA CTTCGACGGA GACGGGATGC TGGACCTCAT CTTGTCCCAT 960 GGAGAGTCCA TGGCTCAGCC GCTGTCCGTC TTCCGGGGCA ATCAGGGCTT CAACAACAAC
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PCT/US02/12476

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Protein Accession #: Eos sequence

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	OATOTOTO O	74444444444444444444444444444444444444	TTTACTACTG	AGTCAAGTTT	TCTAGTTCTC	TGTAATTGTT	7680
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WO 02/086443 Coding sequence: 1..714

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WO 02/086443
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PCT/US02/12476

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          CTCCTGCTGG TGATCTGCCT GATCTTCTCC ATCATGGGCT ACTACTATGT TCCTGTAAAG
                                                                                             2100
          ACAGAGGATA TGCGGGGTCC AGCAGATAAG CACATTCCTC ACATCCAGGG GAACATGATC
                                                                                            2160
           AAACTAGAGA CCAAGAAGAC AAAACTCTGA
  85
           Seg ID NO: 681 Protein sequence
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Protein Accession #: AAB34388.1

PCT/US02/12476

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WO 02/086443
                                  21
                                               31
                                                            41
                                                                         51
                     11
        MNPFOKNESK ETLPSPVSIE EVPPRPPSPP KKPSPTICGS NYPLSIAFIV VNEFCERFSY
        YGMKAVLILY FLYFLHWNED TSTSIYHAFS SLCYFTPILG AAIADSWLGK FKTIIYLSLV
                                                                                        120
 5
        YVLGHVIKSL GALPILGGQV VHTVLSLIGL SLIALGTGGI KPCVAAFGGD QFEEKHABER
        TRYFSVFYLS INAGSLISTF ITPMLRGDVQ CFGEDCYALA FGVPGLLMVI ALVVFAMGSK
                                                                                         240
        IYNKPPPEGN IVAQVFKCIW FAISNRFKNR SGDIPKRQHW LDWAAEKYPK QLIMDVKALT
                                                                                        300
        RVLFLYIPLP MFWALLDQQG SRWTLQAIRM NRNLGFFVLQ PDQMQVLNPF LVLIFIPLFD
                                                                                        360
        RVIFILIES HEMAINSLER AVGMILACIA FAVABAVEIK INEMAPAGS POEVFLQVIN
LADDEVKVIV VCNENNSLLI ESIKSFQKTP HYSKLHLKTK SQDPHFHLKY HNLSLYTEHS
10
                                                                                         480
        VQEKNWYSLV IREDGNSISS MMVKDTESKT TNGMTTVRFV NTLHKDVNIS LSTDTSLNVG
                                                                                        540
        EDYGYSAYRT VORGEYPAVH CRTEDKNFSL NLGLLDFGAA YLFVITNNYN QGLQAMKIED
IPANKMSIAW QLPQYALVTA GEVMFSVTGL EFSYSQAPSS MKSVLQAAWL LTIAVGNIIV
                                                                                         600
                                                                                         660
        LVVAQFSGLV QWAEFILFSC LLLVICLIFS IMGYYYVPVK TEDMRGPADK HIPHIQGNMI
                                                                                        720
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        Nucleic Acid Accession #: NM_016077.1
        Coding sequence: 128..667
20
                     11
        TOGCTTTGTG ATTCTTGATC CGGAACTTTG TCACCCAGGA ACCCCGGAAG AGGTAGCTCA
        25
        ACTGTAGATG CCCTCCAAAT CCTTGGTTAT GGAATATTTG GCTCATCCCA GTACACTCGG
                                                                                         180
        CTTGGCTGTT GGAGTTGCTT GTGGCATGTG CCTGGGCTGG AGCCTTCGAG TATGCTTTGG
                                                                                         240
        GATGCTCCC AAAAGCAAGA CAGACAAGAC ACACACAGAT ACTGAAAGTG AAGCAAGCAT
CTTGGGAGAC AGCGGGGAGT ACAAGATGAT TCTTGTGGTT CGAAATGACT TAAAGATGG
                                                                                         360
        AAAAGGGAAA GTGGCTGCCC AGTGCTCTCA TGCTGCTGTT TCAGCCTACA AGCAGATTCA
                                                                                         420
        AAGAAGAAAT CCTGAAATGC TCAAACAATG GGAATACTGT GGCCAGCCCA AGGTGGTGGT
30
                                                                                         480
        CAAAGCTCCT GATGAAGAAA CCCTGATTGC ATTATTGGCC CATGCAAAAA TGCTGGGACT
        GACTGTAAGT TTAATTCAAG ATGCTGGACG TACTCAGATT GCACCAGGCT CTCAAACTGT
                                                                                         600
        CCTAGGGATT GGGCCAGGAC CAGCAGACCT AATTGACAAA GTCACTGGTC ACCTAAAACT TTACTAGGTG GACTTTGATA TGACAACAAC CCCTCCATCA CAAGTGTTTG AAGCCTGTCA GATTCTTAACA ACAAAAGCTG AATTTCTTCA CCCAACTTAA ATGTTCTTGA GATGAAAATA
                                                                                         660
                                                                                         780
35
        AAACCTATTC CCATGTTCTA AAAAAA
        Seg ID NO: 683 Protein seguence
        Protein Accession #: NP_057161.1
40
        MPSKSLVMBY LAHPSTLGLA VGVACGMCLG WSLRVCFGML PKSKTSKTHT DTESEASILG
                                                                                          60
        DSGEYKMILV VRNDLKMGKG KVAAQCSHAA VSAYKQIQRR NPEMLKQWEY CGQPKVVVKA
45
        PDEETLIALL AHAKMLGLTV SLIQDAGRTQ IAPGSQTVLG IGPGPADLID KVTGHLKLY
                      684 DNA sequence
        Nucleic Acid Accession #: NM_004864.1
        Coding sequence: 26..952
50
                                  21
        CGGAACGAGG GCAACCTGCA CAGCCATGCC CGGGCAAGAA CTCAGGACGG TGAATGGCTC
TCAGATGCTC CTGGTGTTGC TGGTGCTCTC GTGGCTGCCG CATGGGGGCG CCCTGTCTCT
55
        GGCCGAGGCG AGCCGCGCAA GTTTCCCGGG ACCCTCAGAG TTGCACTCCG AAGACTCCAG
                                                                                         180
        ATTCCGAGAG TTGCGGAAAC GCTACGAGGA CCTGCTAACC AGGCTGCGGG CCAACCAGAG
CTGGGAAGAT TCGAACACCG ACCTCGTCCC GGCCCCTGCA GTCCGGATAC TCACGCCAGA
                                                                                         240
        AGTGCGGCTG GGATCCGGCG GCCACCTGCA CCTGCGTATC TCTCGGGCCG CCCTTCCCGA
                                                                                         360
        GGGGCTCCCC GAGGCCTCCC GCCTTCACCG GGCTCTGTTC CGGCTGTCCC CGACGGCGTC
                                                                                         420
        AGGETCGTG GAGGTGACAC GACCGCTGCG GCGTCAGCT AGCCTTGCAA GACCCCAAGC
GCCCGCGCTG CACCTGCGAC TGTCGCCGCC GCCGTCGCAG TCGGACCAAC TGCTGGCAGA
60
                                                                                         540
        ATCTTCGTCC GCACGGCCCC AGCTGGAGTT GCACTTGCGG CCGCAAGCCG CCAGGGGGCG
                                                                                         600
        CCGCAGAGCG CGTGCGCCCA ACGGGGACGA CTGTCCGCTC GGGCCCGGGC GTTGCTGCCG
TCTGCACACG GTCCGCGCGT CGCTGGAAGA CCTGGGCTGG GCCGATTGGG TGCTGTCGCC
                                                                                         660
                                                                                         720
65
        ACGGGAGGTG CAAGTGACCA TGTGCATCGG CGCGTGCCCG AGCCAGTTCC GGGCGGCAAA
                                                                                         780
        CATGCACGGG CAGATCAAGA CGAGCCTGCA CCGCCTGAAG CCCGACACGG AGCCAGCGCC
CTGCTGCTG CCCGCCAGCT ACAATCCCAT GGTGCTCATT CAAAAGACCG ACACCGGGGT
                                                                                         840
        GTCGCTCCAG ACCTATGATG ACTTGTTAGC CAAAGACTGC CACTGCATAT GAGCAGTCCT
                                                                                         960
        GGTCCTTCCA CTGTGCACCT GCGCGGGGGA GGCGACCTCA GTTGTCCTGC CCTGTGGAAT
                                                                                        1020
        GGGCTCAAGG TTCCTGAGAC ACCCGATTCC TGCCCAAACA GCTGTATTTA TATAAGTCTG
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                                                                                        1080
        TTATTTATTA TTAATTTATT GGGGTGACCT TCTTGGGGAC TCGGGGGCTG GTCTGATGGA
        ACTGTGTATT TATTTAAAAC TCTGGTGATA AAAATAAAGC TGTCTGAACT GTTAAAAAAA
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        Seq ID NO: 685 Protein sequence
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        MPGQELRTVN GSQMLLVLLV LSWLPHGGAL SLAEASRASF PGPSELHSED SRFRELRKRY
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                                                                                          60
        EDLLTRLRAN QSWEDSNTDL VPAPAVRILT PEVRLGSGGH LHLRISRAAL PEGLPEASRL
                                                                                         120
         HRALFRLSPT ASRSWDVTRP LRRQLSLARP QAPALHLRLS PPPSQSDQLL AESSSARPQL
                                                                                         180
         ELHLRPQAAR GRRRARARNG DDCPLGPGRC CRLHTVRASL EDLGWADWVL SPREVQVTMC
                                                                                         240
         IGACPSQFRA ANMHAQIKTS LHRLKPDTEP APCCVPASYN PMVLIQKTDT GVSLQTYDDL
85
         LAKDCHCI
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Seq ID NO: 686 DNA sequence

Nucleic Acid Accession #: NM_002423.2 Coding sequence: 48..851

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                                                                                  120
       GAGGCATGAG TGAGCTACAG TGGGAACAGG CTCAGGACTA TCTCAAGAGA TTTTATCTCT
                                                                                  180
       ATGACTCAGA AACAAAAAT GCCAACAGTT TAGAAGCCAA ACTCAAGGAG ATGCAAAAAT
                                                                                  240
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       TCTTTGGCCT ACCTATAACT GGAATGTTAA ACTCCCGCGT CATAGAAATA ATGCAGAAGC
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        CCAGATGTGG AGTGCCAGAT GTTGCAGAAT ACTCACTATT TCCAAATAGC CCAAAATGGA
                                                                                  360
        CTTCCAAAGT GGTCACCTAC AGGATCGTAT CATATACTCG AGACTTACCG CATATTACAG
                                                                                  420
        TGGATCGATT AGTGTCAAAG GCTTTAAACA TGTGGGGCAA AGAGATCCCC CTGCATTTCA
                                                                                  480
       GGAAAGTTGT ATGGGGAACT GCTGACATCA TGATTGGCTT TGCGCGAGGA GCTCATGGGG
                                                                                  540
15
       ACTCCTACCC ATTTGATGGG CCAGGAAACA CGCTGGCTCA TGCCTTTGCG CCTGGGACAG
                                                                                  600
       GTCTCGGAGG AGATGCTCAC TTCGATGAGG ATGAACGCTG GACGGATGGT AGCAGTCTAG
                                                                                  660
       GGATTAACTT CCTGTATGCT GCAACTCATG AACTTGGCCA TTCTTTGGGT ATGGGACATT
                                                                                  720
       CCTCTGATCC TAATGCAGTG ATGTATCCAA CCTATGGAAA TGGAGATCCC CAAAATTTTA
                                                                                  780
       AACTITCCCA GGATGATATI AAAGGCATIC AGAAACTATA TGGAAAGAGA AGTAATICAA
                                                                                  840
20
       GAAAGAAATA GAAACTTCAG GCAGAACATC CATTCATTCA TTCATTGGAT TGTATATCAT
                                                                                  900
       TGTTGCACAA TCAGAATTGA TAAGCACTGT TCCTCCACTC CATTTAGCAA TTATGTCACC
                                                                                  960
        CTTTTTTATT GCAGTTGGTT TTTGAATGTC TTTCACTCCT TTTATTGGTT AAACTCCTTT
                                                                                 1020
       ATGGTGTGAC TGTGTCTTAT TCCATCTATG AGCTTTGTCA GTGCGCGTAG ATGTCAATAA
       ATGTTACATA CACAAATAAA TAAAATGTTT ATTCCATGGT AAATTTA
25
       Seq ID NO: 687 Protein sequence
Protein Accession #: NP_002414.1
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                                            31
                                                        41
                                                                   51
                    11
30
       MRLTVLCAVC LLPGSLALPL PQEAGGMSEL QWEQAQDYLK RPYLYDSETK NANSLEAKLK
                                                                                   60
       EMQKFFGLPI TGMLNSRVIE IMQKPRCGVP DVAEYSLFPN SPKWTSKVVT YRIVSYTRDL
PHITVDRLVS KALNMWGKEI PLHFRKVVWG TADIMIGFAR GAHGDSYPFD GPGNTLAHAF
                                                                                  120
                                                                                  180
       APGTGLGGDA HFDEDERWTD GSSLGINFLY AATHELGHSL GMGHSSDPNA VMYPTYGNGD
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        PONFKLSODD IKGIQKLYGK RSNSRKK
       Seg ID NO:
                    688 DNA sequence
       Nucleic Acid Accession #: NM_005221.3
       Coding sequence: 1..870
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       ATGACAGGAG TGTTTGACAG AAGGGTCCCC AGCATCCGAT CCGGCGACTT CCAAGCTCCG
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       TTCCAGACGT CCGCAGCTAT GCACCATCCG TCTCAGGAAT CGCCAACTTT GCCCGAGTCT
45
       TCAGCTACCG ATTCTGACTA CTACAGCCCT ACGGGGGGAG CCCCGCACGG CTACTGCTCT
                                                                                  180
        CCTACCTCGG CTTCCTATGG CAAAGCTCTC AACCCCTACC AGTATCAGTA TCACGGCGTG
                                                                                  240
       AACGGCTCCG CCGGGAGCTA CCCAGCCAAA GCTTATGCCG ACTATAGCTA CGCTAGCTCC
                                                                                  300
       TACCACCAGT ACGGCGGCGC CTACAACCGC GTCCCAAGCG CCACCAACCA GCCAGAGAAA
                                                                                  360
       GAAGTGACCG AGCCCGAGGT GAGAATGGTG AATGGCAAAC CAAAGAAAGT TCGTAAACCC
                                                                                  420
50
       AGGACTATT ATTCCAGCTT TCAGCTGGCC GCATTACAGA GAAGGTTTCA GAAGACTCAG
TACCTCGCCT TGCCGGAACG CGCCGAGCTG GCCGCCTCGC TGGGATTGAC ACAAACACAG
                                                                                  480
                                                                                  540
       GTGAAAATCT GGTTTCAGAA CAAAAGATCC AAGATCAAGA AGATCATGAA AAACGGGAG
ATGCCCCCGG AGCACAGTCC CAGCTCCAGC GACCCAATGG CGTGTAACTC GCCGCAGTCT
                                                                                  600
                                                                                  660
       CCAGCGGTGT GGGAGCCCCA GGGCTCGTCC CGCTCGCTCA GCCACCACCC TCATGCCCAC
                                                                                  720
55
       CCTCCGACCT CCAACCAGTC CCCAGCGTCC AGCTACCTGG AGAACTCTGC ATCCTGGTAC
                                                                                  780
       ACAAGTGCAG CCAGCTCAAT CAATTCCCAC CTGCCGCCGC CGGGCTCCTT ACAGCACCCG
                                                                                  840
       CTGGCGCTGG CCTCCGGGAC ACTCTATTAG
       Seq ID NO: 689 Protein sequence
60
       Protein Accession #: NP_005212.1
                                            31
                                                        41
                                                                   51
                               21
       MTGVFDRRVP SIRSGDFQAP FQTSAAMHHP SQESPTLPES SATDSDYYSP TGGAPHGYCS
                                                                                   60
65
       PTSASYGKAL NPYQYQYHGV NGSAGSYPAK AYADYSYASS YHQYGGAYNR VPSATNQPEK
                                                                                  120
       EVTEPEVRMV NGKPKKVRKP RTIYSSFQLA ALQRRFQKTQ YLALPERAEL AASLGLTQTQ
       VKIWFONKRS KIKKIMKNGE MPPEHSPSSS DPMACNSPQS PAVWEPQGSS RSLSHHPHAH
        PPTSNQSPAS SYLENSASWY TSAASSINSH LPPPGSLQHP LALASGTLY
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It is understood that the examples described above in no way serve to limit the true scope of this invention, but rather are presented for illustrative purposes. All publications, sequences of accession numbers, and patent applications cited in this specification are herein neorporated by reference as if each individual publication or patent application were specifically and individually indicated to be incorporated by reference.

WHAT IS CLAIMED IS:

1	1.	A method of detecting a lung cancer-associated transcript in a cell
2	from a patient, the	method comprising contacting a biological sample from the patient with a
3	polynucleotide tha	at selectively hybridizes to a sequence at least 80% identical to a sequence
4	as shown in Table	s 1A-16.
1	2.	The method of claim 1, wherein the polynucleotide selectively
2	hybridizes to a sec	quence at least 95% identical to a sequence as shown in Tables 1A-16.
1	3.	The method of claim 1, wherein the biological sample is a tissue
2	sample.	
1	4.	The method of claim 1, wherein the biological sample comprises
2	isolated nucleic ac	
1	5.	The method of claim 4, wherein the nucleic acids are mRNA.
1	6.	The method of claim 4, further comprising the step of amplifying
2	nucleic acids before	re the step of contacting the biological sample with the polynucleotide.
1	7.	The method of claim 1, wherein the polynucleotide comprises a
2	sequence as shown	n in Tables 1A-16.
1	8.	The method of claim 1, wherein the polynucleotide is labeled.
1	9.	The method of claim 8, wherein the label is a fluorescent label.
1	10.	The method of claim 1, wherein the polynucleotide is immobilized on
2	a solid surface.	
1	11.	The method of claim 1, wherein the patient is undergoing a therapeutic
2	regimen to treat lu	ng cancer.
1	. 12.	The method of claim 1, wherein the patient is suspected of having lung
2	cancer.	
1	13.	A method of monitoring the efficacy of a therapeutic treatment of lung

cancer, the method comprising the steps of:

2

PCT/US02/12476 WO 02/086443 3 (i) providing a biological sample from a patient undergoing the therapeutic 4 treatment; and 5 (ii) determining the level of a lung cancer-associated transcript in the 6 biological sample by contacting the biological sample with a polynucleotide that selectively 7 hybridizes to a sequence at least 80% identical to a sequence as shown in Tables 1A-16, 8 thereby monitoring the efficacy of the therapy. 1 14. The method of claim 13, further comprising the step of: (iii) comparing 2 the level of the lung cancer-associated transcript to a level of the lung cancer-associated transcript in a biological sample from the patient prior to, or earlier in, the therapeutic 3 4 treatment. 1 15. The method of claim 13, wherein the patient is a human. 1 16. A method of monitoring the efficacy of a therapeutic treatment of lung 2 cancer, the method comprising the steps of: 3 (i) providing a biological sample from a patient undergoing the therapeutic 4 treatment; and 5 (ii) determining the level of a lung cancer-associated antibody in the biological 6 sample by contacting the biological sample with a polypeptide encoded by a polynucleotide 7 that selectively hybridizes to a sequence at least 80% identical to a sequence as shown in 8 Tables 1A-16, wherein the polypeptide specifically binds to the lung cancer-associated 9 antibody, thereby monitoring the efficacy of the therapy. 1 17. The method of claim 16, further comprising the step of: (iii) comparing 2 the level of the lung cancer-associated antibody to a level of the lung cancer-associated 3 antibody in a biological sample from the patient prior to, or earlier in, the therapeutic 4 treatment. 1 18. The method of claim 16, wherein the patient is a human. 1 19. A method of monitoring the efficacy of a therapeutic treatment of lung 2 cancer, the method comprising the steps of: 3 (i) providing a biological sample from a patient undergoing the therapeutic

4

treatment; and

5	(ii) determining the level of a lung cancer-associated polypeptide in the		
6	biological sample by contacting the biological sample with an antibody, wherein the antibod		
7	specifically binds to a polypeptide encoded by a polynucleotide that selectively hybridizes to		
8	a sequence at least 8	0% identical to a sequence as shown in Tables 1A-16, thereby	
9	monitoring the effic	acy of the therapy.	
1	20.	The method of claim 19, further comprising the step of: (iii) comparing	
1		cancer-associated polypeptide to a level of the lung cancer-associated	
2	_	logical sample from the patient prior to, or earlier in, the therapeutic	
3		logical sample from the patient prior to, of earlier in, the morapoune	
4	treatment.		
1	21.	The method of claim 19, wherein the patient is a human.	
1	22.	An isolated nucleic acid molecule consisting of a polynucleotide	
2	sequence as shown i	in Tables 1A-16.	
	-		
1	23.	The nucleic acid molecule of claim 22, which is labeled.	
1	24.	The nucleic acid of claim 23, wherein the label is a fluorescent label	
1	25.	An expression vector comprising the nucleic acid of claim 22.	
1	26.	A host cell comprising the expression vector of claim 25.	
1	27.	An isolated polypeptide which is encoded by a nucleic acid molecule	
2	having polynucleoti	de sequence as shown in Tables 1A-16.	
1	28.	An antibody that specifically binds a polypeptide of claim 27.	
1	29.	The antibody of claim 28, further conjugated to an effector component.	
1	30.	The antibody of claim 29, wherein the effector component is a	
2	fluorescent label.		
1	31.	The antibody of claim 29, wherein the effector component is a	
2	radioisotope or a cy	totoxic chemical.	
1	. 20	The entitledy of claim 20, which is an antibody fragment	

1	33	The antibody of claim 29, which	Il is a numainzed antibody
1	34	A method of detecting a lung of	ancer cell in a biological sample from a
2	patient, the metho	omprising contacting the biologic	cal sample with an antibody of claim
3	28.		
1	35	The method of claim 34, where	ein the antibody is further conjugated to
2	an effector compo		
1	26	The method of claim 35, where	ein the effector component is a
1 2	36 fluorescent label.	The method of claim 33, where	em the effector component is a
1	37	_	ies specific to lung cancer in a patient,
2	-	_	e from the patient with a polypeptide
3	encoded by a nuc	acid comprises a sequence from	Tables 1A-16.
1	38	A method for identifying a cor	npound that modulates a lung cancer-
2	associated polype	e, the method comprising the st	eps of:
3	(i)	tacting the compound with a lur	ng cancer-associated polypeptide, the
4	polypeptide enco	by a polynucleotide that selective	vely hybridizes to a sequence at least
5		uence as shown in Tables 1A-1	·
6	(ii	ermining the functional effect of	of the compound upon the polypeptide.
1	39	The method of claim 38, where	ein the functional effect is a physical
2	effect.		
1	40	The method of claim 38, when	ein the functional effect is a chemical
2	effect.	,	
1	41	The method of claim 38, when	ein the polypeptide is expressed in a
	eukaryotic host c		the projection of the projecti
2	eukaryouc nost c	r cen memorane.	
1	42	The method of claim 38, where	ein the functional effect is determined by
2	measuring ligand	ding to the polypeptide.	
1	43	The method of claim 38, where	ein the polypeptide is recombinant.

1	44.	A method of inhibiting proliferation of a lung cancer-associated cell to	
2	treat lung cancer in a patient, the method comprising the step of administering to the subject		
3	therapeutically effective amount of a compound identified using the method of claim 38.		
1	45.	The method of claim 44, wherein the compound is an antibody.	
1	46.	The method of claim 45, wherein the patient is a human.	
1	47.	A drug screening assay comprising the steps of	
2	(i) ad	ministering a test compound to a mammal having lung cancer or a cell	
3	isolated therefrom;		
4	(ii) co	omparing the level of gene expression of a polynucleotide that selectively	
5	hybridizes to a sequence at least 80% identical to a sequence as shown in Tables 1A-16 in a		
6	treated cell or mamr	nal with the level of gene expression of the polynucleotide in a control	
7	cell or mammal, wh	erein a test compound that modulates the level of expression of the	
8	polynucleotide is a	candidate for the treatment of lung cancer.	
1	48.	The assay of claim 47, wherein the control is a mammal with lung	
2	cancer or a cell there	efrom that has not been treated with the test compound.	
1	49.	The assay of claim 47, wherein the control is a normal cell or mammal	
1	50.	A method for treating a mammal having lung cancer comprising	
2	administering a com	pound identified by the assay of claim 47.	
1	51.	A pharmaceutiPcal composition for treating a mammal having lung	
2	cancer, the composi	tion comprising a compound identified by the assay of claim 47 and a	
3	physiologically acceptable excipient.		

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of inventorship (Rule 4.17(iv)) for US only

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PATENT COOPERATION TREATY

PCT

DECLARATION OF NON-ESTABLISHMENT OF INTERNATIONAL SEARCH REPORT

(PCT Article 17(2)(a), Rule 13ter.1(c) and 39)

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Applicant's or agent's file reference 18501-15-3PC	IMPORTANT DECL	ARATION	Date of mailing (day/month/year) 15 AUG 2003	
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Applicant		-		
EOS BIOTECHNOLOGY, INC			•	
a. scientific theories. b. mathematical theories. c. plant varieties. d. animal varieties. e. essential biological and the products of schemes, rules or methods for treatments. j. methods for treatments. k. diagnostic methods	application for the reasons indicate an application relates to: des processes for the production of a such processes, nethods of doing business, nethods of performing purely menthods of playing games, ent of the human body by surger practised on the human or animal practised on the human or animal practised on the human or animal practised on the human or animal practised on the human or animal practised on the human or animal practised on the human or animal practised on the human or animal practised on the human or animal practised on the human or animal practised on the human or animal practised on the human or animal practised on the human or animal practised on the human or animal practised on the human or animal practised on the human or animal practised on the human or animal practice.	plants and animals, ntal acts. y or therapy.	o International search report other than microbiological processes	
2. The failure of the following parts of the international application to comply with prescribed requirements prevents a meaningful search from being carried out: the description the claims the drawings				
The failure of the nucleotide and/or amino acid sequence listing to comply with the standard provided for in Annex C of the Administrative Instructions prevents a meaningful search from being carried out: the written form has not been furnished or does not comply with the standard. the computer readable form has not been furnished or does not comply with the standard.				
4. Further comments:				
Name and mailing address of the ISA/US Mail Stop PCT, Atm: ISA/US Commissioner for Patents P.O. Box 1450 Alexandria, Virginia 22313-1450 Parsimile No. (703)305-3230 Authorized officer Carla Myers Carla Myers PRIMARY EXAMINER Telephone No. 703-308-0196				
Pacsimile No. (703)305-3230 Form PCT/ISA/203 (July 1998)		Zotophone 110. 70	5-503-0120	

PATENT COOPERATION TREATY

To: TOWNSEND AND TOWNSEND AND CREW LLP TWO EMBARCADERO CENTER EIGHTH FLOOR SAN FRANCISCO, CA 94111-3834	PCT NOTIFICATION OF TRANSMITTAL OF THE INTERNATIONAL SEARCH REPORT OR THE DECLARATION (PCT Rule 44.1) Date of Mailing (day/month/year) 15 AUG 2003			
Applicant's or agent's file reference 18501-15-3PC	FOR FURTHER ACTION See paragraphs 1 and 4 below			
International application No. PCT/US92/12476	International filing date (day/month/year) 18 April 2002 (18.04.2002)			
Applicant EOS BIOTECHNOLOGY, INC	1			
1. The applicant is hereby notified that the international search report has been established and is transmitted herewith. Filing of amendments and statement under Article 19: The applicant is entitled, if he so wishes, to amend the claims of the international application (see Rule 46): When? The time limit for filing such amendments is normally two months from the date of transmittal of the international search report. Where? Directly to the International Bureau of WIPO, 34, chemin des Colombettes 1211 Geneva 20, Switzerland, Facsimile No.: (41-22) 740.14.35 For more detailed instructions, see the notes on the accompanying sheet.				
2. The applicant is hereby notified that no international search report will be established and that the declaration under Article 17(2)(a) to that effect is transmitted herewith.				
3. With regard to the protest against payment of (an) additional fee(s) under Rule 40.2, the applicant is notified that: the protest together with the decision thereon has been transmitted to the International Bureau together with the applicant's request to forward the texts of both the protest and the decision thereon to the designated Offices. no decision has been made yet on the protest; the applicant will be notified as soon as a decision is made.				
4. Reminders Shortly after 18 months from the priority date, the international application will be published by the International Bureau. If the applicant wishes to avoid or postpone publication, a notice of withdrawal of the international application, or of the priority claim, must reach the International Bureau as provided in Rules 90 bis.1 and 90 bis.3, respectively, before the completion of the technical preparations for international publication. Within 19 months from the priority date, but only in respect of some designated Offices, a demand for international preliminary				
examination must be filed if the applicant wishes to postpone the entry into the national phase until 30 months from the priority date (in some Offices even later); otherwise the applicant must, within 20 months from the priority date, perform the prescribed acts for entry into the national phase before those designated Offices. In respect of other designated Offices, the time limit of 30 months (or later) will apply even if no demand is filed within 19 months.				
See the Annex to Form PCT/IB/301 and, for details about the applicable time limits, Office by Office, see the PCT Applicant's Guids, Volume II, National Chapters and the WIPO Internet site.				
Name and mailing address of the ISA/US Mail Stop PCT, Attn: ISA/US Commissioner for Patents P.O. Box 1450 Alexandria, Virginia 22313-1450 Facsimile No. (703)305-3230 Form PCT/ISA/220 (April 2002) Authorized officer Carlie Myels Carlie Myels Carlie Myels Carlie Myels Carlie Myels (See notes on accompanying sheet)				